Characterization of *Epichloë coenophiala* within the US: are all tall fescue endophytes created equal?

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Tall fescue (*Lolium arundinaceum*) is a valuable and broadly adapted forage grass that occupies approximately 14 million hectares across the United States. A native to Europe, tall fescue was likely introduced into the US around the late 1800’s. Much of the success of tall fescue can be attributed to *Epichloë coenophiala* (formerly *Neotyphodium coenophialum*) a seed borne symbiont that aids in host persistence. *Epichloë* species are capable of producing a range of alkaloids (ergot alkaloids, indole-diterpenes, lolines, and peramine) that provide protection to the plant host from herbivory. Unfortunately, most tall fescue within the US, commonly referred to as “Kentucky-31” (KY31), harbors the endophyte *E. coenophiala* that causes toxicity to grazing livestock due to the production of ergot alkaloids. Molecular analyses of tall fescue endophytes have identified four independent associations, representing tall fescue with *E. coenophiala*, *Epichloë* sp. FaTG-2, *Epichloë* sp. FaTG-3, or *Epichloë* sp. FaTG-4. Each of these *Epichloë* species can be further distinguished based on genetic variation that equates to differences in the alkaloid gene loci. Tall fescue samples were evaluated using markers to simple sequence repeats (SSRs) and alkaloid biosynthesis genes to determine endophyte strain variation present within continental US. Samples represented seed and tillers from the Suiter farm (Menifee County, KY), which is considered the originating site of KY31, as well as plant samples collected from 14 states, breeder’s seed and plant introduction lines (National Plant Germplasm System, NPGS). This study revealed two prominent *E. coenophiala* genotypes based on presence of alkaloid biosynthesis genes and SSR markers and provides insight into endophyte variation within continental US across historical and current tall fescue samples.

**Keywords:** *Neotyphodium coenophialum*, fescue toxicosis, ergot alkaloids, Kentucky-31 (KY31), endophyte diversity

**INTRODUCTION**

**HISTORY OF KY31 TALL FESCUE AND DISCOVERY OF ENDOPHYTE**

Tall fescue [*Lolium arundinaceum* (Schreb.) Darbysh. syn Festuca arundinacea Schreb.] was introduced into the United States from Europe in the late 1800’s and is considered an important cool season perennial forage crop (Hoveland, 2009). Tall fescue is widely adapted to the eastern United States spanning 14 million hectares (35 million acres) with the fescue belt considered the major region of adaptation and use (Figure 1) (Ball et al., 1993). A timeline representing significant research events of tall fescue is shown in Figure 1 and outlined below.

One of the most well known tall fescue cultivars, “Kentucky-31” (KY31), was collected in 1931 by Dr. E. N. Fergus (University of Kentucky) on a farm owned by William Suiter (Menifee County, KY) (Fergus and Buckner, 1972). KY31 gained wide acceptance as a grass with excellent agronomic attributes under difficult growth conditions, such as drought and poor soils. The KY31 ecotype was released in 1942, but was not officially registered as a cultivar until 1972 (Fergus and Buckner, 1972). Subsequently, the persistence and success of KY31 was attributed to the presence of the systemic fungal endophyte, *Epichloë coenophiala*. It was estimated that 90% of all tall fescue pastures in the US are endophyte infected (Siegel et al., 1985). The fitness benefits the endophyte provides include drought tolerance, improved competitive ability (Arachevaleta et al., 1989; West et al., 1993; Malinowski and Belesky, 2000), as well as protection from herbivores through the production of bioactive alkaloids (Clay et al., 1985; Bacon et al., 1986).

Unfortunately, although KY31 was known as a persistent cultivar, reports of poor animal performance when grazing this forage began in the 1940s (Cunningham, 1949; Jacobson et al., 1963). Animals that grazed on tall fescue suffered maladies such as fescue foot, fat necrosis, and fescue toxicosis (Bush et al., 1979). Cattle experiencing fescue toxicosis can exhibit rough hair coats, heat stress (wallowing in mud), elevated rectal temperatures, vasoconstriction, suppressed appetite, reduced prolactin levels, poor growth (lower average daily gains), and a reduction in calving rates (Hoveland et al., 1983; Hemken et al., 1984;
TALL FESCUE ENDOPHYTE DESCRIPTION

Initially, Bacon et al. (1977) identified the tall fescue endophyte as *E. typhina*, which was later renamed *Acremonium coenophialum* to acknowledge the anamorphic state of *Epichloë* species (Morgan-Jones and Gams, 1982). Christensen and Latch (1991) described variation among isolates of *A. coenophialum* from tall fescue, and in 1993 the taxonomy of these endophytes was described (Christensen et al., 1993). The genus *Acremonium* was reclassified using phylogenetic analyses and *A. coenophialum* was renamed *Neotyphodium coenophialum* (Glenn et al., 1996). Finally, under the nomenclatural rule changes for fungi *Neotyphodium* is now included in the genus *Epichloë* resulting in the change to *E. coenophiala* (Leuchtmann et al., 2014). *E. coenophiala* is considered an asexual hybrid, phylogenetically described as a triparental hybrid with inferred ancestral progenitors from *E. festucae*, *E. typhina* subsp. *poeae*, and the *Lolium*-associated endophyte (LAE) (*Tsai et al., 1994; Moon et al., 1999*).

*E. coenophiala*, like many *Epichloë* species, is capable of producing a variety of bioactive secondary metabolites. The four described classes of alkaloids produced by *Epichloë* species are ergot alkaloids, indole-diterpenes, lolines, and peramine (Siegel et al., 1990). Ergot alkaloids (e.g., ergovaline) and the indole-diterpene, lolitrem B, have been shown to have anti-mammalian activity causing fescue toxicosis (Bacon et al., 1977) and rye-grass staggers (Fletcher and Harvey, 1981), respectively. Peramine is considered an insect feeding deterrent (Johnson et al., 1985; Rowan and Latch, 1994) and the lolines have been documented for their potent insecticidal activity (Bush et al., 1997). *E. coenophiala* as a species complex can produce all four classes of alkaloids (Table 1). However, the most commonly produced alkaloids are peramine, lolines, and ergovaline (Christensen et al., 1993; Leuchtmann et al., 2000; Schardl et al., 2013b).

Considerable research has been conducted to understand the biosynthesis of these bioactive compounds including identification and characterization of the gene products required for the biosynthesis of each alkaloid class (Panaccione et al., 2001; Wang et al., 2004; Spiering et al., 2005, 2008; Tanaka et al., 2005; Young et al., 2006, 2009; Fleetwood et al., 2007; Saikia et al., 2012; Pan et al., 2014). This has been supported with genome sequences, including draft genome sequences of three *E. coenophiala* strains (Schardl et al., 2013a,b). This research has provided an understanding of why *Epichloë* species can have diverse alkaloid profiles and provided the sequence to develop markers for mating type and key alkaloid biosynthesis genes to genetically evaluate endophyte diversity *in planta* (Charlton et al., 2012, 2014; Takach et al., 2012; Takach and Young, 2014).

To date, tall fescue is known to form associations with four taxonomic groups, *E. coenophiala*, *Epichloë* sp. FaTG-2, *Epichloë* sp. FaTG-3, and *Epichloë* sp. FaTG-4 that vary based on ploidy (either 2x or 3x) and progenitors (Table 1). Initially tall fescue endophytes were distinguished by morphology and isozyme analysis to establish taxonomic groupings, and variation was also seen with the production of peramine, ergovaline, lolitrem B, and lolines (Christensen et al., 1993). Phylogenetic analyses were able
Table 1 | *Epichloë* species and genotype variation associated with endophytes of tall fescue.

| Endophyte species | Endophyte genotype<sup>a</sup> | Ploidy | Progenitors<sup>b</sup> | Minimum marker set to distinguish species genotypes | Predicted chemotype class<sup>d</sup> |
|------------------|-----------------------------|-------|----------------------|---------------------------------|------------------|
| *Epichloë coenophiala* | Profile 1 | 3x | Efe (II) × LAE (Vb) × Ety (Ib) | AAA | dmaW, lpsB, idtG, idtQ, lolC, lolA, perA-A2 | EAS, LOL, PER |
| *Epichloë coenophiala* | Profile 2 | 3x | Efe (II) × LAE (Vb) × Ety (Ib) | AAA | dmaW, lpsB, idtQ, lolC, lolA, perA-A2 | EAS, LOL, PER |
| *Epichloë coenophiala* | Profile 3 | 3x | Efe (II) × LAE (Vb) × Ety (Ib) | AAA | dmaW, lpsB, idtG, idtQ, lolC, lolA, perA-A2 | EAS, IDT, LOL, PER |
| *Epichloë coenophiala* | Profile 4 | 3x | Efe (II) × LAE (Vb) × Ety (Ib) | AAA | idtG, idtQ, lolC, lolA, perA-A2 | IDT, LOL, PER |
| *Epichloë* sp. | FaTG-2 | Profile 1 | 2x | Efe (II) × LAE (Vb) | BB | dmaW, lpsB, idtG, idtQ, ltmJ, perA-A2, ΔperA-A2 | EAS, LTM, PER |
| *Epichloë* sp. | FaTG-2 | Profile 2 | 2x | Efe (II) × LAE (Vb) | AB | dmaW, lpsB, idtG, idtQ, perA-A2, ΔperA-A2 | EAS, IDT, PER |
| *Epichloë* sp. | FaTG-3 | Profile 3 | 2x | Efe (II) × LAE (Vb) | AB | dmaW, lpsB, idtG, idtQ, ltmJ, perA-A2, ΔperA-A2 | EAS, LTM, PER |
| *Epichloë* sp. | FaTG-3 | Profile 4 | 2x | LAE (Vb) × Ety (Ia) | AA | idtG, idtQ, lolC, lolA, perA-A2, ΔperA-A2 | IDT, LOL, PER |
| *Epichloë* sp. | FaTG-4 | Profile 5 | 2x | LAE (Vb) × Ety (Ia) | AA | idtG, idtQ, lolC, lolA, perA-A2, ΔperA-A2 | LOL, PER |
| *Epichloë* sp. | FaTG-4 | Profile 6 | 2x | LAE (Vb) × Ety (Ia) | AB | dmaW, lpsB, idtG, idtQ, perA-A2 | EAS, IDT, PER |

<sup>a</sup>Based on designations from Takach and Young (2014) and draft genome sequences of *FaTG-2* isolates NFe45079 and NFe45115.

<sup>b</sup>Efe (II), *E. festucae* (matting population II); LAE, *Lolium* associated endophyte (matting population Vb); Ety (Ia), *E. typhina* (matting population 1a); Ety (Ib), *E. typhina* (matting population 1b); Mating population as designated from Leuchtmann et al. (2014).

<sup>c</sup>The perA-A2 marker is designed to the second adenylation domain. Some isolates have a deletion in this domain as represented by perA-A2 (Takach et al., 2012).

<sup>d</sup>Predicted chemotype class represents the class of genes that are found in the genome and do not always represent a functionally active locus. EAS, ergot alkaloids; IDT, indole diterpenes; LTM, lolitrem B; LOL, lolines; PER, peramine.

to define the relationships of each taxonomic group to distinguish the ancestral progenitors of these hybrid species (Schardl et al., 1991, 2013b; Moon et al., 2004). The most studied of these species is *E. coenophiala*, the endophyte first identified in KY31.

Isozyme analyses of *E. coenophiala* isolates from within the US indicated that very little variation existed within this species (Leuchtman and Clay, 1990). Although isozyme analysis can reflect endophyte diversity, this analysis requires pure cultures and thus is limited by the number of samples per tall fescue line that can be screened. Genetic analysis can now be performed directly with endophyte infected plant material using high throughput systems (Takach and Young, 2014). Markers have recently been used to determine the genetic diversity between tall fescue endophyte isolates and also evaluate their potential for alkaloid production (Ekanayake et al., 2012; Takach et al., 2012; Takach and Young, 2014). In fact, variation of mating type and alkaloid genes determined by PCR could be enough to allow placement of tall fescue endophytes into distinct genotype groups associated with each *Epichloë* species (Table 1) (Takach and Young, 2014). At least four unique *E. coenophiala* genotypes are easily distinguishable among tall fescue sourced originally from Europe and the Mediterranean basin (Ekanayake et al., 2012; Takach and Young, 2014).

Literature surrounding endophyte-infected tall fescue that causes fescue toxicosis often refers to *E. coenophiala* as the common toxic endophyte. The objective of this study was to compare the endophytes within tall fescue cultivars, varieties and ecotypes from the US using markers to SSRs and alkaloid biosynthesis genes to identify and characterize these endophytes. We have determined endophyte diversity across historical and current tall fescue samples to evaluate the endophyte diversity that may exist across the US.

MATERIALS AND METHODS

**BIOLOGICAL MATERIALS**

Tall fescue plant material was provided by researchers in Alabama, Arkansas, Georgia, Kentucky, Missouri, Mississippi, New York, Ohio, Oklahoma,Pennsylvania, South Carolina, Tennessee, Texas, and West Virginia (Table 2). Plants were maintained in a space plant nursery under rain-fed conditions or in the greenhouse at the Samuel Roberts Noble Foundation, Ardmore, Oklahoma. Each researcher was requested to provide at least 10 independent
Table 2 | Sources of tall fescue plants from US collection.

| State       | County          | Plant designationa | No. of plants maintained | No. plants that died | No. of endophyte-infected plants | Endophyte genotypes present |
|-------------|-----------------|--------------------|--------------------------|----------------------|----------------------------------|-------------------------------|
| AL          | Dallas          | Black belt station | 21                       | 0                    | 21                               | 2-1, 2-2                     |
| AR          | Nevada          | Prescott           | 6                        | 0                    | 6                                | 1-1, 2-1                     |
| AR          | Hempstead       | Deanne             | 6                        | 0                    | 6                                | 2-1                          |
| FL, GA, MD, NY | GA-5           | Walker             | 10                       | 2                    | 7                                | 1-1, 1-2, 2-1, 2-3           |
| GA          | Walker          | Walker county      | 12                       | 0                    | 8                                | 2-1                          |
| GA          | Wayne           | Jesup              | 9                        | 2                    | 6                                | 2-1                          |
| KY          | Mennifee        | Suiter farm        | 10                       | 1                    | 9                                | 1-1, 2-1, 2-4                |
| KY          | Caldwell        | Pennyville         | 10                       | 1                    | 4                                | 2-1                          |
| MO          | Camden          | Lake Farm          | 12                       | 0                    | 9                                | 2-1                          |
| MO          | Camden          | Ford place         | 8                        | 0                    | 8                                | 2-1                          |
| MO          | Camden          | Tiny’s place       | 4                        | 0                    | 3                                | 2-1                          |
| MO          | St. Louis       | Hencken            | 6                        | 0                    | 6                                | 2-1                          |
| MS          | Oktibbeha       | Starksville        | 9                        | 0                    | 9                                | 2-1, 2-4                     |
| NY          | Allegany        | Alfred             | 5                        | 0                    | 5                                | 2-1                          |
| OH          | Coshocton       | NAEW graze         | 6                        | 0                    | 6                                | 1-1, 2-1                     |
| OH          | Coshocton       | NAEW hay           | 6                        | 0                    | 5                                | 2-1                          |
| OK          | Carter          | NFTF 1000          | 9                        | 0                    | 7                                | 2-1                          |
| OK          | Woodward        | NFTF 1100          | 9                        | 1                    | 7                                | 2-1                          |
| OK          | Hughes          | Calvin             | 2                        | 0                    | 1                                | 2-1                          |
| PA          | Huntingdon      | Soder              | 1                        | 0                    | 1                                | 1-7                          |
| PA          | Huntingdon      | Soder (Petersburg) | 3                        | 0                    | 3                                | 3-1                          |
| PA          | Centre          | Everhart           | 1                        | 0                    | 1                                | 2-3                          |
| PA          | Centre          | JRE state college  | 5                        | 1                    | 4                                | 2-1, 2-2                     |
| SC          | Anderson        | NFTF 1491          | 10                       | 0                    | 5                                | 2-1                          |
| TN          | Henderson       | Lexington          | 4                        | 0                    | 4                                | 2-1                          |
| TN          | Henderson       | Natchez trace      | 7                        | 0                    | 6                                | 1-1, 2-1                     |
| TX          | Fannin          | Fannin             | 4                        | 0                    | 4                                | 2-1                          |
| TX          | Fannin          | NFTF 1492          | 10                       | 3                    | 6                                | 2-1, 2-2                     |
| TX          | Crosby, Lubbock | NFTF 1230          | 9                        | 0                    | 8                                | 1-1, 2-1                     |
| TX          | Kerr            | NFTF 1480          | 9                        | 1                    | 8                                | 2-1                          |
| WV          | Raleigh         | Roscoe upper       | 10                       | 0                    | 8                                | 1-1, 2-1                     |
| WV          | Raleigh         | Roscoe middle      | 10                       | 1                    | 7                                | 2-1                          |
| WV          | Raleigh         | Roscoe lower       | 10                       | 3                    | 6                                | 2-1, 2-4                     |
| WV          | Raleigh         | Reba               | 10                       | 3                    | 5                                | 2-1                          |

aPlant designation refers to landmark or site location or plant breeding line information (NFTF).

plants from fields known to cause fescue toxicosis or thought to contain the common toxic endophyte. Tall fescue seed stocks (PI lines) were sourced from the National Plant Germplasm System (NPGS). Georgia-5 (GA-5) seed was provided by JH Bouton and other seed stocks were sourced from the Samuel Roberts Noble Foundation tall fescue (NFTF) breeding program and designated NFTF.

DNA ISOLATION AND ENDOPHYTE GENOTYPING

Total DNA from individual seeds or tillers from stock plants were isolated using QIAGEN MagAttract 96 DNA Plant Core Kit (Qiagen Inc., Valencia, CA). Primers specific for tefA, tef1-exon1d (5′-GGGTAAGGACGAAAAGACTCA-3′) and tef1-exon5u-1 (5′-CGGCACCGATAATCAGGATAG-3′) (Craven et al., 2001; Moon et al., 2002) were used to detect the presence of endophyte. A minimum set of key alkaloid genes and two mating type genes were chosen to differentiate the E. coenophiala endophytes present in continental tall fescue based on the previous study by Takach and Young (2014). The markers were designed to mtAC and mtBA mating type genes, dmaW and lpsB for representatives of the EAS locus, lolC and lolA for representatives of the LOL locus, idtG and idtQ for representatives of the IDT locus, and perA second adenylation domain (perA-A2) for PER. Multiplex PCR was performed in a total volume of 25 µL containing 3 µL DNA, 1.0 U GoTaq™ DNA Polymerase (Promega Corp., Madison, WI), 1× Green GoTaq™ Reaction Buffer containing 1.5 mM MgCl₂, 0.2 mM of each dNTP (Promega Corp.), and 1 µM of each primer as described previously (Takach et al., 2012; Charlton et al., 2014; Takach and Young, 2014). The cycling parameters were an initial denaturation step for 1 min at 94 C, 30 cycles of denaturation at
94°C for 15 s, annealing at 56°C for 30 s, extension at 72°C for 45 s, followed by a final synthesis step at 72°C for 10 min.

PCR of the microsatellite B10 and B11 loci (Moon et al., 1999) were used to differentiate endophytes within an E. coenophiala profile. For SSR analysis, one primer at each locus was end labeled with a fluorescent phosphoramidite dye. Specifically, primers B10.1 was labeled with 2′-chloro-7′-phenyl-1,4-dichloro-6-carboxy-fluorescein (VIC) and B11.1 was labeled with 2′-chloro-5′-fluoro-7′,8′-benzo-1,4-dichloro-6-carbo xyfluorescein (NED) (Life Technologies, Carlsbad, CA). PCR was performed in a total volume of 10 μL containing diluted DNA (approximately 0.5 ng), 0.75 U Platinum Taq DNA Polymerase (Life Technologies), 1 × PCR Buffer (-Mg), 1.5 mM MgCl₂, 100 nM of each dNTP (Promega Corp.) and 200 nM of each primer. The cycling parameters were an initial denaturation step for 4 min at 94°C, 35 cycles of denaturation at 94°C for 30 s, annealing at 60°C for 30 s, extension at 72°C for 30 s, followed by a final synthesis step at 72°C for 7 min. PCR products (1.5 μL of a 1:10 dilution) were added to 9.9 μL of Hi-Di formamide and 0.1 μL of GeneScan™ 500 LIZ™ size standard (Life Technologies). Samples were denatured at 94°C for 5 min prior to separation on an ABI 3730 DNA Analyzer. Data analysis was performed using Peak Scanner Software v1.0 (Applied Biosystems).

ERGOVALINE ANALYSIS

Pseudostems were collected from greenhouse grown plants, lyophilized and ground into a fine powder and stored at −20°C. Ergovaline concentrations were measured in duplicate using 10 µL of methanol containing 0.005 mg/mL dihydroergotamine tartrate salt (Sigma-Aldrich, St. Louis, MO) for 3 h and then centrifuged at 1800 × g for 5 min. Each sample was analyzed for the presence of ergovaline using an ACQUITY ultra-performance liquid chromatography (UPLC) system (Waters Corporation, Milford, MA) as described previously (Takahashi et al., 2012). Seed extracts in which the ergovaline content was previously quantified (A. M. Craig, Endophyte Testing Laboratory, Oregon State University) were used as standards for quantification. Concentrations used to generate the standard curve included 0, 50, 97, 500, 1000, and 2000 ppb. The linear standard curve was plotted as ergovaline:ergotamine peak area ratio vs. the actual amount of ergovaline. Standards and samples were analyzed in duplicate and their values averaged.

RESULTS AND DISCUSSION

EVALUATION OF HISTORICAL TALL FESCUE ENDOPHYTES FROM THE UNITED STATES

The cultivar KY31 is well known for causing fescue toxicosis and has been distributed over much of the eastern United States (Figure 1). KY31 was established from an ecotype collection from the Suiter farm in Menifee County, KY (released as a cultivar in 1943) and source material (PI 531431) from this location was deposited into NPGS in 1991. Other cultivars, developed after KY31, have also contributed to the dissemination of endophyte-infected tall fescue (Pedersen and Sleper, 1988). In particular, Alta (cultivar in 1945) was considered a successful cultivar in northeast US and was likely interbred with KY31 (Asay et al., 1979). Seed from other early tall fescue cultivars such as Alta, Kenmont (cultivar in 1963), Kenwell (cultivar in 1965), Kenhy (cultivar in 1977), and Missouri 96 (cultivar in 1977) (Pedersen and Sleper, 1988) were also included in our study (see Table 3 for NPGS deposition

Table 3 | Characterization of endophytes from Kentucky 31 tall fescue seed in US based on microsatellite variation.

| Seed stock        | Year | Seeds tested | %E+ (number) | %E− (number) | %E+ Ecotype profile | %E− Ecotype profile |
|-------------------|------|--------------|--------------|--------------|---------------------|---------------------|
| PI 561431 - KY31  | 1991 | 46           | 91% (42)     | 9% (4)       | 48% (22)            | 43% (20)            |
| KY31 commercial seed | 2011 | 48         | 79% (38)     | 21% (10)     | 17% (8)             | 63% (30)            |
| KY31 - SW Missouri | 2008 | 46         | 98% (45)     | 2% (1)       | 0% (0)              | 98% (45)            |
| PI 596701 - Missouri 96 | 1979 | 24         | 0            | 100 (24)    | 0% (0)              | 0% (0)              |
| PI 578714 - Kenmont | 1963 | 24         | 0            | 100 (24)    | 0% (0)              | 0% (0)              |
| PI 574521 - Kenwell | 1965 | 22         | 0            | 100 (22)    | 0% (0)              | 0% (0)              |
| PI 434051 - Kenhy | 1979 | 24         | 0            | 100 (24)    | 0% (0)              | 0% (0)              |
| PI 601020 - Johnstone | 1983 | 24         | 0            | 100 (24)    | 0% (0)              | 0% (0)              |
| PI 578712 - Alta | 1962 | 48         | 0            | 100 (48)    | 0% (0)              | 0% (0)              |
| Jesup E+ | 2003 | 10         | 100% (10)    | 0% (0)       | 0% (0)              | 100% (10)           |
| NFTF 1000 - PDF E+ | 1998 | 36         | 100% (36)    | 0% (0)       | 8% (3)              | 92% (33)            |
| NFTF 1011 - PDF E+ | 2007 | 12         | 100% (12)    | 0% (0)       | 33% (4)             | 67% (8)             |
| NFTF 1041 - PDF E+ | 2010 | 12         | 100% (12)    | 0% (0)       | 0% (0)              | 100% (12)           |

*aThe year the seed was purchased, generated or when it entered NPGS.

*bAs determined by markers consistent with E. coenophiala profile 1 and 2 from Table 1.

*cSSR B10 = 161, 170, 184; B11 = 147, 191.

*dSSR B10 = 152, 161, 178; B11 = 171, 195.

*eThe KY31 commercial seed Tri-Star Seed Co., Inc. located in Spring Hill, KS was purchased from the Tractor Supply Company, Ardmore, Oklahoma, in July 2011.

+fJohnstone was released as an endophyte free cultivar (Buckner et al., 1983).

*gNFTF 1041 and NFTF 1041 represent selections from the original NFTF 1000 (also known as PDF E+ Hopkins et al., 2011) from Oklahoma.
dates). Studies that have included some of these early cultivars have subsequently indicated they were infected with a common toxic endophyte (Bacon et al., 1977; Cornell et al., 1982; Siegel et al., 1984; Pedersen and Sleper, 1988). We also evaluated more recent cultivars and germplasm from the NFTF breeding program that contain common toxic endophyte (Table 3).

Seed from each cultivar or line were analyzed for endophyte infection and genetic variation (Table 3). Unfortunately many of the seed samples sourced from NPGS were endophyte-free or had levels less than 5%. Since endophyte viability can be compromised during storage (Siegel et al., 1985; Rolston and Agee, 2007) there was no guarantee these seeds would represent the endophyte status of the original plant material. Only the three KY31 samples, Jesup E+ and NFTF breeding lines were positive for endophyte presence. In addition, two endophyte genotypes, E. coenophiala profile 1 and profile 2 that vary based on presence of IDT genes, could be distinguished in four of these seed samples. However, the percentage of each endophyte strain varied in each seed lot. Four of the tested seed lines, KY31 (from Missouri), Jesup E+ and NFTF 1041 may represent an E. coenophiala profile 2 monoculture, or contain a low incidence of E. coenophiala profile 1 as the number of seeds tested for some lines were low (Table 3).

Lines that were selected from NFTF 1000-PDF (NFTF 1011 selected from PDF for vigorous growth, high forage yield and digestibility, and NFTF 1041 selected from PDF for high digestibility by marker assisted selection) showed different ratios of each E. coenophiala profile (Table 3). This may show the influence an endophyte strain, which is maintained in the maternal line, can have on selectable traits such as persistence and vigor if the endophyte provides a host advantage. As such, it would be interesting to evaluate the host genetic shifts under selection, with and without endophytes, while also following selection of different endophyte strains.

ENDOPHYTE ANALYSIS FROM DEVELOPMENT OF THE CULTIVAR GEORGIA-5

Analysis of endophyte variation within a population provides an opportunity to evaluate material incorporated through a tall fescue breeding pipeline and eventually released for commercial production (Figure 2). The GA-5 cultivar was developed as a synthetic endophyte-infected cultivar with superior forage yield and persistence in the Southern Coastal Plains that had potential to replace KY31 (Bouton et al., 1993b). The cultivar was established from five clones and was shown to be 75% endophyte infected (Bouton et al., 1993b). We evaluated seed from the original five clones (each clone having originated from a different location) using markers to SSRs and alkaloid biosynthesis genes to determine the initial infection rates of each clone and identify which E. coenophiala profiles were present. The endophyte status of the originating lines varied from 32 to 100% infection, and the endophyte profiles were consistent within the seed sample from each clone. Three independent endophyte genotypes (based on SSRs) were identified within the clones (Figure 2). Seed from synthetic 1 established in 1980 was also tested for endophyte infection and identification, and all three endophyte genotypes were represented within this sample with an overall endophyte infection level of 79% (Figure 2). In 1993, GA-5 was registered as a cultivar (Bouton et al., 1993b) and subsequently released commercially in 1996. When we evaluated a seed stock from the commercial line the overall infection level was 69% and two of the three expected endophyte SSR profiles were identified within the sample. However, an additional endophyte genotype (B10 = 152, 161, 178 and B11 = 171, 195) was present in 5% of the seed sample (Figure 2) that has likely arisen from contamination later in production. The level of endophyte free seed increased from Syn 1 (21%) to Syn 6 (31%) and may indicate that production favored this part of the population. Unfortunately we were unable to detect the endophyte genotype profile 2 with B10 = 161, 173, 178 and B11 = 171, 195, which may be due to the number of seeds that were tested.

ENDOPHYTE DIVERSITY FROM US TALL FESCUE COLLECTIONS

To further examine tall fescue endophyte diversity within the US, collections of tall fescue plants from 14 eastern states were evaluated for endophyte presence. In total, 246 tall fescue plants were screened using markers for SSRs B10 and B11, and the minimum set of alkaloid biosynthesis genes to distinguish the different E. coenophiala genetic profiles. Of the 246 plants initially screened, 204 (83%) were endophyte-infected. All of the endophyte-infected samples could be amplified with primer sets to mtAC, perA, and the LOL and EAS markers, and samples only varied with the presence of IDT markers. Three E. coenophiala genetic profiles were identified (E. coenophiala profiles 1, 2, and 3; Table 4).

Further analysis using the B10 and B11 SSR markers showed additional variation within the genetic profiles. There were two unique SSR profiles for genotype 1, four SSR profiles for genotype 2 and one SSR profile for genotype 3. E. coenophiala profile 1 is consistent with the same pattern of the KY31 endophyte E. coenophiala strain e19 (Schardl et al., 1991; Takach and Young, 2014), while E. coenophiala profile 2 is more highly represented in the samples.

The two most common SSR marker patterns were also identified in the KY31 seed (Table 3). Interestingly, of the nine plants that were collected from the original Suiter farm (Table 2), one plant showed a third unique SSR profile in common with endophytes found in West Virginia and Mississippi. Only one set of plants from Pennsylvania produced E. coenophiala genetic profile 3, but unfortunately knowledge about the tall fescue planted at this site was unavailable. This Pennsylvania endophyte-infected tall fescue is the only set of plants that were likely to produce indole-diterpenes.

Plants with unique E. coenophiala profiles were selected from the nursery and maintained in the greenhouse. Ergovaline was analyzed from pseudostems of 25 plants representing the seven unique E. coenophiala profiles. Each of the plant-endophyte associations produced ergovaline (Table 5), indicating that the endophyte in these lines would all be considered a common toxic endophyte. The ergovaline levels generated under greenhouse conditions were also extremely high, well above the threshold of 400–750 ppb required for toxicity in livestock (Hovermale and Craig, 2001). Although the levels tested from these plants are high, it may simply reflect they were well maintained plants subjected to regular fertilization, which is known to result
in higher levels of ergovaline production (Rottinghaus et al., 1991).

This study provides a snapshot of common toxic endophyte-infected tall fescue across the eastern US. The endophyte genotypes we identified were consistent with samples from Europe where tall fescue was originally sourced (Takah and Young, 2014). Endophyte genotype diversity was present at the origin of KY31 but it appears *Epichloë coenophiala* genotype 2 is the dominant endophyte genotype across the US. This endophyte was also prevalent in other breeding populations such as Jesup E+ and NFTF 1000. Given the sample numbers we have selected from each location, it is still possible that other endophyte genotypes are present at low frequencies. Although KY31 is believed to be the predominant source of tall fescue in the US, other sources were developed and planted at various times and locations, and it is possible that during this process one endophyte genotype provided a selectable advantage. Indeed, if we look at NFTF 1011, a selection from the NFTF 1000 line (Table 3), we see a shift in endophyte genotype that may be due to specific selection pressures.

**NEW CULTIVARS WITH SELECTED ENDOPHYTES**

To overcome fescue toxicosis, researchers initially removed the endophyte from toxic tall fescue. Unfortunately, studies that evaluated the endophyte role on tall fescue performance found endophyte-infected lines had better persistence and greater yield than endophyte-free lines (Pedersen et al., 1990; Bouton et al., 1993a, 2002; West et al., 1993; Gunter and Beck, 2004). As expected, when production of ergot alkaloids was low or not present, livestock also had better overall performance and
increased average daily gains (Stuedemann and Hoveland, 1988; Gunter and Beck, 2004). An ideal solution to capture both endophyte associated plant persistence and reduced livestock toxicity was to identify an endophyte strain that retained traits for plant persistence, but did not produce the alkaloids toxic to livestock. Analysis of tall fescue across its natural distribution of Europe and the Mediterranean basin has shown diversity of both the endophyte (Christensen et al., 1993; Ekanayake et al., 2012; Takach and Young, 2014) and its plant host (Hand et al., 2012); subsequently this diversity has been exploited to establish selected endophyte-infected tall fescue with low mammalian toxicity (reviewed in Bouton, 2009; Johnson et al., 2013; Young et al., 2013). Typically these endophytes lack many or all of the genes at the EAS locus required for ergot alkaloid biosynthesis (e.g., *E. coenophiala* profile 4, Table 1) (Takach and Young, 2014), although some endophytes have been selected for lower ergot alkaloid production.

Initial success with selected endophytes was observed when the endophyte strain AR542 (known commercially as MaxQ and MaxP in the US and Australia, respectively) was inoculated into Jesup MaxQ (Pennington Seed, Inc.) was the first commercial tall fescue cultivar to be released and used by farmers containing a selected endophyte. Additional endophyte-infected tall fescue lines have since been established and evaluated for both plant and animal performance (Roberts and Andrae, 2004; Hopkins et al., 2010; Parish et al., 2013; Beck et al., 2014). One of the most recently released cultivars, Texoma MaxQ II, is the result of breeding for plant persistence with an ecotypic selection that was subsequently inoculated with the endophyte strain AR584 (MaxQ II) that does not cause livestock toxicity (Hopkins et al., 2010, 2011). Current commercially available cultivars of selected endophyte-tall fescue associations include: Jesup MaxQ (Pennington Seed, Inc.), Texoma MaxQ (Pennington Seed Inc.), BarOptima Plus E34 (Barenbrug), Duramax Gold (DLF International Seeds), and Estancia with ArkShield (MountainView Seeds).

### Table 4 | Characterization of endophytes from tall fescue plants in US based on alkaloid profiles and microsatellite variation.

| Endophyte genotype | B10 allele sizes (bp) | B11 allele sizes (bp) | PER<sup>a</sup> | EAS<sup>b</sup> | LOL<sup>c</sup> | IDT<sup>d</sup> | No. of plants | States |
|--------------------|-----------------------|-----------------------|----------------|----------------|----------------|----------------|---------------|--------|
| *E. coenophiala* profile 1-1 | 161, 170, 184 | 147, 191 | + | + | + | + | 11 | AR, KY, OH, TN, TX, WV |
| *E. coenophiala* profile 1-2 | 139, 170, 184 | 163, 191 | + | + | + | + | 2 | GA |
| *E. coenophiala* profile 2-1 | 152, 161, 178 | 171, 195 | + | + | + | + | 172 | AL, AR, GA, KY, MO, MS, NY, OH, OK, PA, SC, TN, TX, WV |
| *E. coenophiala* profile 2-2 | 152, 161, 178 | 183, 195 | + | + | + | + | 3 | AL, TX |
| *E. coenophiala* profile 2-3 | 161, 173, 178 | 171, 195 | + | + | + | + | 3 | GA (NY)<sup>b</sup>, PA |
| *E. coenophiala* profile 2-4 | 161, 173, 178 | 171, 210 | + | + | + | + | 4 | KY, MS, WV |
| *E. coenophiala* profile 3-1 | 161, 170, 178 | 155, 163 | + | + | + | + | 3 | PA |

<sup>a</sup>The + indicates that a PCR fragment was detected with primers designed to the gene.

<sup>b</sup>Identified out of GA-5 and represents the clone from NY (see Figure 2).

### Table 5 | Ergovaline concentrations of tall fescue infected with different *Epichloë coenophiala* genotypes.

| Endophyte genotype<sup>a</sup> | B10 allele sizes (bp) | B11 allele sizes (bp) | No. of plants tested | States represented | Range of [Ergovaline] (ppm) |
|------------------------------|-----------------------|-----------------------|----------------------|-------------------|-----------------------------|
| *E. coenophiala* profile 1-1 | 161, 170, 184 | 147, 191 | 5 | GA, KY, TN, TX, WV | 2.2–7.1 |
| *E. coenophiala* profile 1-2 | 139, 170, 184 | 163, 191 | 1 | GA | 4.1 |
| *E. coenophiala* profile 2-1 | 152, 161, 178 | 171, 195 | 11 | AL, GA, KY, MO, MS, PA, TN, TX, WV | 1.7–7.8 |
| *E. coenophiala* profile 2-2 | 152, 161, 178 | 183, 195 | 2 | AL, TX | 1.2–9.3 |
| *E. coenophiala* profile 2-3 | 161, 173, 178 | 171, 195 | 3 | GA (NY)<sup>b</sup>, PA | 2.4–5.5 |
| *E. coenophiala* profile 2-4 | 161, 173, 178 | 171, 210 | 2 | KY, MS | 1.9–8.4 |
| *E. coenophiala* profile 3-1 | 161, 170, 178 | 155, 163 | 1 | PA | 4.2 |

<sup>a</sup>Based on designations from Takach and Young (2014).

<sup>b</sup>Identified from the GA-5 line and represents the clone from NY (see Figure 2).

### CONCLUSION

Endophytes have clearly affected the success of tall fescue within the US, from the prevalence of the common toxic endophyte through to advancing cultivars with selected endophytes. Many farmers have learned to manage the effects of fescue toxicity through pasture management techniques, but now there are also options for eliminating toxicity by pasture replacement. The
availability of elite tall fescue lines infected with selected endophytes allows farmers to provide nutritious, non-toxic feed for their livestock without fear of toxic repercussions. Climate change will likely increase the dependency of tall fescue monocultures to rely upon endophytes to provide drought tolerance and protection from insect pests. As more selected endophyte-infected cultivars enter the marketplace to replace tall fescue pastures containing common toxic endophytes, we will be able to examine the constancy of these symbiotic associations over time. Utilization of molecular markers will provide effective methods to identify endophyte strains within tall fescue cultivars, varieties and ecotypes, and help distinguish endophyte friend from foe.

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