Different nitrogen acquisition patterns of plant and soil microorganisms in the forestgrassland transition zone on the Loess Plateau

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Figure. S1 Locations of the study region and four studied stands in the forest–grassland transition zone on the Loess Plateau, China.



Figure. S2 Rainfall (mm) and 0–50cm soil moisture content (%) in the four studied stands before and after ¹⁵N labeling. Arrows represent the rainfall and moisture content at 3, 7, and 15 days after labeling.



Time



Figure. S3 The average proportion of root surface area between 0–20 cm, 20–50 cm, 50–100 cm in the range of 0–100 cm in the four studied stands (means \pm SE, n = 9).

Figure. S4 The δ^{15} N of different plant organs in the tree stands of (a, b) *H. rhamnoides* and (c, d) *P. tabuliformis* (means ± SE, n = 3). Asterisks indicate significant differences in the ¹⁵N enrichment between N forms within each stand (ANOVA and repeated measurements. *, *P* < 0.05; **, *P* < 0.01). D: Time since ¹⁵N labeling (day); L: The form of applied ¹⁵N–labeled N. (p) and (m) represent pure and mixed stands, respectively.



Figure. S5 Uptake rates of ¹⁵N tracer of leaves, branches, fine roots collected from 0–20 cm, 20–20 cm, and 0–50 cm soil in the tree stands of (a, b) *H. rhamnoides* and (c, d) *P. tabuliformis* (means \pm SE, n = 3). Asterisks indicate significant differences in the ¹⁵N uptake rates between N forms within each stand (ANOVA and repeated measurements. *, *P* < 0.05; **, *P* < 0.01). D: Time since ¹⁵N labeling (day); L: The form of applied ¹⁵N–labeled N. (p) and (m) represent pure and mixed stands, respectively.



Figure. S6 The δ^{15} N and uptake rates of ¹⁵N tracer of leaves (a, c) and fine roots (b, d) in *A*. *gmelinii* stand (means ± SE, n = 3). Asterisks indicate significant differences in the ¹⁵N uptake rates between N forms within each stand (ANOVA and repeated measurements. *, *P* < 0.05; **, *P* < 0.01). D: Time since ¹⁵N labeling (day); L: The form of applied ¹⁵N–labeled N.



Figure. S7 The relative proportion of ${}^{15}NH_4^+$ and ${}^{15}NO_3^-$ to total ${}^{15}N$ tracer uptake by the whole plant in the four stands of (a, d) *H. rhamnoides*, (b, e) *P. tabuliformis* and (c) *A. gmelinii* at 3, 7 and 15 days after ${}^{15}N$ labeling, respectively (shown are N–pool–weighted mean by roots, leaves and branches, n = 3). (p) and (m) represent pure and mixed stands, respectively.



Figure. S8 The relative proportion of ${}^{15}NH_4^+$ and ${}^{15}NO_3^-$ to total ${}^{15}N$ tracer uptake by different plant organs (roots, leaves and branches) and whole plant (N–pool–weighted mean) in four studied stands of (a, d) *H. rhamnoides*, (b, e) *P. tabuliformis* and (c) *A. gmelinii* at 3, 7 and 15 days after ${}^{15}N$ labeling (n = 3). (p) and (m) represent pure and mixed stands, respectively.



Figure. S9 The relative proportion of ${}^{15}NH_4^+$ and ${}^{15}NO_3^-$ to total ${}^{15}N$ tracer uptake by soil microorganisms in the four stands at 3, 7 and 15 days after ${}^{15}N$ labeling, respectively (n = 3).

