Introduction of H₂-Uptake Hydrogenase Genes Into Rhizobial Strains Improves Symbiotic Nitrogen Fixation in *Vicia sativa* and *Lotus corniculatus* Forage Legumes

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Biological nitrogen fixation by the *Rhizobium*-legume symbiosis allows the conversion of atmospheric nitrogen into ammonia within root nodules mediated by the nitrogenase enzyme. Nitrogenase activity results in the evolution of hydrogen as a result of a side reaction intrinsic to the activity of this enzyme. Some rhizobia, and also other nitrogen fixers, induce a NiFe uptake hydrogenase (Hup) to recycle hydrogen produced by nitrogenase, thus improving the efficiency of the nitrogen fixation process. In this work we report the generation and symbiotic behavior of hydrogenase-positive *Rhizobium leguminosarum* and *Mesorhizobium loti* strains effective in vetch (*Vicia sativa*) and birsfoot trefoil (*Lotus corniculatus*) forage crops, respectively. The ability of hydrogen recycling was transferred to these strains through the incorporation of *hup* minitransposon TnHB100, thus leading to full recycling of hydrogen in nodules. Inoculation of *Vicia* and *Lotus* plants with these engineered strains led to significant increases in the levels of nitrogen incorporated into the host legumes. The level of improvement of symbiotic performance was dependent on the recipient strain and also on the legume host. These results indicate that hydrogen recycling has the potential to improve symbiotic nitrogen fixation in forage plants.

Keywords: *hup* genes, *Rhizobium leguminosarum*, *Mesorhizobium loti*, legume, inoculant

INTRODUCTION

Symbiotic nitrogen fixation, carried out by rhizobia expressing nitrogenase in association with legume plants, allows reducing the use of N fertilizers, thus positioning legume crops as key players in sustainable agriculture (Laranjo et al., 2014). Nitrogen fertilizers used in agriculture are incorporated into plants with low efficiency (40–50%) resulting in the release of an enormous amount of reactive nitrogen into the biosphere (Zhang et al., 2020). Forage legumes (alfalfa, clover, vetch, trefoil, etc.) can be used as cut fodder (either directly or after fermentation) or as grazed pasture, and are also relevant at a global scale for ruminant’s feeding.
Common vetch (Vicia sativa) is a multi-purpose, cool season annual legume widely used in low- to medium rainfall areas as source of both forage and grain for animal feeding, due to its high protein content (Huang et al., 2017). V. sativa has a high potential of nitrogen fixation through its association with endosymbiotic bacteria. V. rhizobia were traditionally placed within Rhizobium leguminosarum species, but systematic analysis of isolates from different regions has revealed that other rhizobial species such as R. sophorae and R. laguerrae, are also endosymbionts of this legume (Zhang et al., 2019).

Lotus is a widely spread legume genus comprising 100–130 species originally native from Europe, Asia, Africa and Australia. Its current worldwide distribution is a consequence of human-driven introduction into non-native areas due to its interest as forage legume and its adaptability to different environmental stresses, which make it a relevant alternative in ecological restoration programs (Escaray et al., 2012). Just a few species, including Lotus corniculatus, L. subbiflorus, and L. tenuis, have been domesticated and subject to plant breeding programs for its use as forage for livestock. Among them, L. corniculatus is a major component of pastures in South America (Díaz et al., 2005). Lotus rhizobia are widely diverse, with over 20 species dispersed amongst several genera (Mesorhizobium, Bradyrhizobium, Rhizobium, Ensifer and Aminobacter; Lorite et al., 2018). The study of the Lotus symbiosis has been greatly advanced by the establishment of L. japonicus as a model system for genetic and molecular studies (Tabata and Stougaard, 2014).

Growth of pastures based in forage legumes is closely associated with the rate of biological N fixation, since nitrogen is often the main limiting nutrient for herbage production (Phelan et al., 2015). Nitrogenase activity results in the evolution of hydrogen as a result of a side reaction of the enzyme, thus accounting for at least 25% of electrons channeled into the enzyme. As a consequence, nodules in root legumes evolve large amounts of hydrogen, which diffuses into the soil, constituting a source of inefficiency of the system; some rhizobial strains induce a hydrogenase system able to recycle the hydrogen produced by nitrogenase, thus improving the efficiency of the nitrogen fixation process (Ruiz-Argüeso et al., 2000; Palacios et al., 2005). This hydrogenase is a metalloenzyme with a complex NiFe heterobimetallic cofactor. As with other metalloenzymes such as nitrogenase, the synthesis of the NiFe hydrogenase is a complex process that, in the case of R. leguminosarum bv. viciae, requires 18 proteins whose genetic determinants (hupSLCDEFGHJKhypABFCDEX) are clustered in the symbiotic plasmid of this bacterium (Palacios et al., 2005). Synthesis of hydrogenase is co-regulated with nitrogenase though the action of NifA, the master regulator of nitrogen fixation genes (Brito et al., 1997).

The cloning of the whole cluster of hydrogenase genes a cosmid (Levya et al., 1987) allowed the introduction of hydrogen-uptake (Hup) activity into hydrogen-deficient strains of different rhizobial species, including R. leguminosarum, R. etli, and Mesorhizobium loti (Brito et al., 2000). In order to avoid instability of the introduced genes derived from its presence as an extrachromosomal entity, a mini-transposon (TnHB100) including the whole set of hup/hyp genes along with a spectinomycin resistance marker gene was generated (Bascones et al., 2000). The use of this transposon allowed the generation of transposants inducing high levels of symbiotic hydrogenase in Hup− recipient strains from B. japonicum, R. etli, and R. leguminosarum by viciae, leading to nodules that evolved little or no hydrogen at all. In contrast, transposants obtained from Sinorhizobium meliloti and M. cicer carrying the same transposon expressed very low levels of hydrogenase activity, resulting in nodules evolving high levels of hydrogen (Bascones et al., 2000).

The evaluation of the effect of hydrogen recycling has been carried out in several systems, and positive effects on the level of nitrogen fixation had been shown in different legumes such as cowpea, and common beans (Baginski et al., 2005; Torres et al., 2020). These experiments had been carried out with grain legumes, and there are scarce data available on the effect of hydrogen recycling on forage legumes. Rhizobial strains effective in alfalfa and in clover lack hydrogenase activity. Strategies for heterologous expression of hup genes, either through plasmid transfer or TnHB100 transposon insertion, have been attempted without much success. The model S. meliloti / alfalfa system does not express hydrogenase under symbiotic conditions, even though vegetative cells of the same Sinorhizobium strain can express high levels of hydrogenase activity (Brito et al., 2002), so it was important to elucidate whether the same system can also improve the growth of forage legumes in which hydrogenase is expressed. In this work we have incorporated the R. leguminosarum hydrogenase genes into rhizobial strains effective on two forage legumes permissive for hydrogenase expression (Lotus and Vicia), and the effect of hydrogenase activity on symbiotic performance of the new strains has been determined.

MATERIALS AND METHODS

Bacterial Strains, Plasmids, and Growth Conditions

R. leguminosarum Rlv13 and Rlv21 were isolated from V. sativa root nodules at Zaidín Estacion Experimental (Granada, Spain). M. loti U261 (=NZP2037, Irisarri et al., 1996) was obtained from the Department of Scientific and Industrial Research, Palmerston South, New Zealand. Rhizobium and Mesorhizobium strains were grown at 28°C in tryptone-yeast extract (TY), yeast-manitol broth (YMB), or Rhizobium minimal media (Rm) (Bascones et al., 2000). Escherichia coli strains were grown at 37°C in Luria-Bertani medium supplemented with the corresponding antibiotics. Minitransposon TnHB100, carrying the entire R. leguminosarum hup cluster was transferred into (Meso)Rhizobium strains by conjugation as previously described (Bascones et al., 2000). For bacterial conjugations, fresh cell cultures of E. coli S17.1 pir(pTnHB100) donor (Simon et al., 1983) and rhizobial recipient strains were mixed on TY plates and incubated overnight at 28°C. Transconjugants were selected on Rm plates supplemented with 200 µg/ml of spectinomycin.

Plant Material and Growth Conditions

Seeds of L. corniculatus cv. La Estanzuela, L. tenuis, L. subbiflorus, and L. uliginosus were surface sterilized with 10%
sodium hypochlorite and pregerminated at 28°C for 2 days in the dark. *V. sativa* cv. Filón seeds were sterilized with ethanol/sodium hypochlorite treatment and pregerminated as previously described (Brito et al., 2000). Three-day old seedlings were sown in pots containing vermiculite supplied with a nitrogen-free nutrient solution (Handberg and Stougaard, 1992; Brito et al., 1994) and inoculated with liquid cultures of rhizobial strains. Plants were maintained in a growth chamber under controlled conditions (16/8 h light/dark; Monza et al., 1992; Brito et al., 2000). Bacteroid suspensions were prepared from freshly detached nodules as previously described (Brito et al., 2008).

**Recombinant DNA Techniques**
Plasmid DNA preparations, restriction enzyme digestions, DNA cloning, transformation of DNA into *E. coli*, agarose gel electrophoresis, Southern hybridizations, and PCR amplifications were performed by standard methods (Sambrook and Russell, 2002). Total DNA from rhizobial strains was isolated as described (Bascones et al., 2000). For the determination of nucleotide sequence flanking the Tn*HB100* insertions, the region was first cloned as a *PstI* restriction fragment in a pBluescript vector plasmid (Agilent Technologies, La Jolla, CA), and the DNA sequence was determined using a primer complementary to the 3′ end of the spectinomycin resistance cassette (5′-GCTGGCTTTCTTGTATCG-3′). Transposon insertions in Rlv13H9 and Rlv21H19 were located in genes encoding potential galactonate dehydratase/epimerases highly similar to protein RLV_6446 from *R. leguminosarum* bv *viciae* (Sanchez-Canizares et al., 2018). These proteins are moderately expressed in pea and lentil bacteroids induced by Rlv UPM791 (Duran et al., 2021), and might function in carbon metabolism pathway, so a potential effect of the gene inactivation on bacterial performance cannot be excluded. In contrast, insertion in Rlv13H6 was located in an intergenic region, with no similarity to described genes, whereas Rlv21H10 carried Tn*HB100* insertion in a hypothetical protein not expressed in bacteroids as deduced from a recent proteomic analysis (Duran et al., 2021). In the case of MI U261H4, the site of insertion was located in an intergenic region downstream of a potassium uptake gene (*kup*) located at positions 2366770-2368683 of the genome (Kelly et al., 2014).

The removal of spectinomycin cassette from Rlv13 and Rlv21 Tn*HB100*-containing derivatives was carried out as described (Bascones et al., 2000). Briefly, DNA regions flanking the resistance marker on each strain were obtained and fused to *hupS* upstream region by restriction cloning in pK18mob8Sac suicide vector (Schäfer et al., 1994). Then, the plasmid was conjugated into the strains carrying the Tn*HB100* insertion, and a double recombination event excising Spc<sup>R</sup> marker was obtained based on the selection provided by *sacB* gene.

**Determination of Nitrogenase and Hydrogenase Activities and Plant Tests**
Nitrogenase activity was determined by the acetylreduction assay, and hydrogenase activity was estimated by measuring bacteroid hydrogen uptake amperometrically, or hydrogen evolution from nodules by gas chromatography (Brito et al., 2000, 2008) using detopped nodulated roots from 35 day-old plants. Plant dry weight was measured after maintaining shoots in an oven at 80°C for 48 h. Nitrogen content of *Lotus* plants was determined from detached leaves collected after 40, 60 and 70 days growing period and analyzed for total nitrogen with a mass spectrometer (IRMS Micromass Isochrom) at SIDI-Laboratorio de Isotopos Estables (Montevideo, Uruguay). Nitrogen content was calculated according with IAEA (IAEA, 2001). Nitrogen content of *Vicia* shoots was determined by a combustion method using a nitrogen analyzer (FP-528-LECO). Experiments were established in a completely random design with at least three replicates. Data were analyzed using ANOVA and means were compared using Fisher’s least significant difference (LSD) test at 5% probability level.

**RESULTS**

**Generation of Hydrogen-Recycling Strains**
In order to analyze the potential effect of hydrogen recycling in symbiotic performance of forage legumes, the *R. leguminosarum* hydrogenase gene cluster was introduced into strains effective in *Vicia sativa* (*R. leguminosarum* bv *viciae* Rlv13 and Rlv21) and in *L. corniculatus* (*Mesorhizobium loti* U261, a broad-range *Lotus* inoculant strain). Stable insertion of *hup* gene package into the recipient strains was carried using transposon Tn*HB100*. Following conjugation with *E. coli* S17.1*pir* (pTn*HB100*), spectinomycin-resistant derivatives arose at frequencies of ca. 10<sup>−4</sup>/recipient (Rlv strains) or 10<sup>−8</sup>/recipient (MI U261). The presence of *hup* genes in spectinomycin-resistant transconjugants was confirmed through identification of *hupS*-hybridizing bands in Southern experiments (Supplementary Figure 1) or through PCR amplification of specific bands corresponding to *hypB* gene using adequate primers (data not shown).

The ability of transposants to effectively recycle hydrogen evolved by nitrogenase was assessed in several derivatives per each recipient strain. In the case of strains effective in *V. sativa*, four derivatives obtained from each of Rlv13 and Rlv21 strains were analyzed for hydrogenase and nitrogenase activities in vetch nodules (Table 1). With one exception, all transconjugants from Rlv strains showed high levels of nitrogenase activity. The exception was the Rlv13 derivative Rlv13H24, which showed a poor symbiotic performance, attributed to a potential deleterious effect of the insertion. Although there were differences in the levels of hydrogenase activity induced in each case, in both sets of strains we found derivatives with levels of hydrogenase activity high enough as to fully recycle the hydrogen evolved by nitrogenase, thus resulting in nodules with no detectable hydrogen evolution: Rlv13H6 and H9, and Rlv21H10 and H19 (Table 1). Insertion sites were determined for these four derivatives (see Materials and Methods), and derivatives Rlv 13H6 and Rlv21H10, bearing insertions with no predicted effect on bacteroid physiology were selected for the analysis of the effect of hydrogen recycling on vetch productivity.

Transposon Tn*HB100* carries a Spc<sup>R</sup> determinant whose presence in bacteria designed to be used as inoculant in the field is not desirable. In order to generate marker-free derivatives more acceptable for field release, strains Rlv13H6 and Rlv21H10 were
TABLE 1 | Nitrogenase and hydrogenase activities of wild-type and engineered strains incorporating hydrogenase genes from *R. leguminosarum* UPM791.

| Strain | Nitrogenase activity$^a$ (molecules H$_2$ mmol protein$^{-1}$ min$^{-1}$) | Hydrogenase activity$^b$ (molecules H$_2$ mmol protein$^{-1}$ min$^{-1}$) | $H_2$ evolution$^c$ (molecules H$_2$ mmol protein$^{-1}$ min$^{-1}$) |
|--------|-------------------------------------------------|-------------------------------------------------|-----------------|
| *R. leguminosarum*/Vicia sativa | | | |
| Rlv13 (WT) | 51.6 ± 2.3 | < 50 | 14.1 ± 0.7 |
| Rlv13H6 | 45.2 ± 5.5 | 1.187 ± 321 | < 0.05 |
| Rlv13H9 | 60.2 ± 0.2 | 1.272 ± 90 | < 0.05 |
| Rlv13H21 | 30.5 ± 1.3 | 0.69 ± 198 | 1.2 ± 0.1 |
| Rlv13H24 | 10 | < 5 | < 0.05 |
| Rlv21 (WT) | 37.9 ± 2.3 | < 50 | 8.7 ± 0.2 |
| Rlv21H10 | 26.8 ± 0.5 | 1.117 ± 546 | < 0.05 |
| Rlv21H16 | 44.1 ± 0.2 | 1.044 ± 506 | 2.5 ± 0.7 |
| Rlv21H19 | 41.4 ± 1.1 | 1.043 ± 473 | < 0.05 |
| Rlv21H22 | 35.4 ± 0.3 | 198 ± 18 | 8.1 ± 0.8 |

*Mesorhizobium loti*/Lotus corniculatus

| Ml U261 (WT) | 2.4 ± 0.8 | n.d. | 1.07 ± 0.4 |
| Ml U261H4 | 0.6 ± 0.2 | n.d. | < 0.05 |
| Ml U261H6 | 1.4 ± 0.3 | n.d. | 0.6 ± 0.1 |
| Ml U261H15 | 1.5 ± 0.1 | n.d. | 0.5 ± 0.3 |
| Ml U261H16 | 1.2 ± 0.1 | n.d. | 0.8 ± 0.1 |
| Ml U261H23 | 1.0 ± 0.1 | n.d. | 0.6 ± 0.3 |
| Ml U261H25 | 2.1 ± 0.7 | n.d. | 0.3 ± 0.1 |

$^a$Values are expressed in μmoles H$_2$ mmol protein$^{-1}$ min$^{-1}$, and represent the mean of at least three determinations ± S.D.

$^b$Values are expressed in nmoles of H$_2$ mmol protein$^{-1}$.

$^c$n.d., not determined.

Further engineered for the removal of SpcR determinant by a double recombination event carried out as described in Materials and Methods. The absence of the spectinomycin resistance cassette in the resulting strains was confirmed by Southern hybridization of genomic DNA using a probe internal to the SpcR gene (Supplementary Figure 1). Both strains, designated as Rlv13H6DS and Rlv21H10DS, induced nodules with levels of nitrogenase activity similar to those in the strains bearing the SpcR determinant. Bacteroids from these strains showed high levels of hydrogenase activity that fully recycled the nitrogenase-derided hydrogen (data not shown).

In the case of *M. loti* U261 derivatives, 6 transconjugants were selected to assess the expression of the introduced hup genes in symbiosis with *L. corniculatus* as the legume host. Each of the transconjugants was used as inocula for *L. corniculatus* plants, and the levels of nitrogenase activity and hydrogen evolution from nodules were determined (Table 1). Levels of nitrogenase activity in the transconjugant strains did not show significant differences with that from the wild type. Nodules induced by the hup-containing derivatives showed significant reductions of the level of hydrogen evolution as compared to nodules induced by the wild-type strain (Table 1). The complete suppression of hydrogen evolution, indicating full recycling of nitrogenase-produced hydrogen, was detected in one of the transconjugants (Ml U261H4). Inspection of the corresponding genomic region (see Materials and Methods section) revealed that this insertion is likely in an intergenic region not affecting the expression of any other gene.

### Analysis of the Effect of Hydrogen Recycling on Plant Productivity

In order to determine the effect of the hydrogen recycling on symbiotic performance of the vetch/*R. leguminosarum* bv viciae system, *V. sativa* plants were inoculated with the wild-type strains Rlv13 and Rlv21 and with the hup-engineered derivatives obtained from them and described above. Inoculated plants were maintained in a growth chamber, and shoot dry matter and nitrogen accumulation were determined 30 days after inoculation. In this analysis, hydrogen-recycling strains derived from Rlv13 showed significant increases ($p < 0.05$) in N fixed per plant (Table 2). Values of shoot dry weight also showed marked increases (24.1% and 21.7 %, respectively) although in this case the higher variability of the data led to a lower significance of the differences ($p < 0.1$). In the case of strain Rlv21, derivatives incorporating the ability of full hydrogen recycling did not show significant variations of any of the two productivity parameters analyzed. These data suggest that the rhizobial background is relevant for the expression of the benefits associated to hydrogen recycling.

Symbiotic performance of *M. loti* U261 and its derivative strain U261H4 were compared in *L. corniculatus* plants grown under controlled conditions. Plant shoot dry weight, and N content of the shoots were determined at 40, 60, and 70 days after inoculation. 60- and 70-day old plants inoculated with the engineered strain showed levels of nitrogen accumulation significantly higher than those inoculated with the wild-type strain. This result was consistently observed in three independent experiments carried out under controlled chamber conditions. The increase in nitrogen was accompanied with an increase in dry weight at 70 days (Table 3). At 40 days plants showed higher values of N accumulation, but the variability of the data made these differences not statistically significant.

Since, *M. loti* U261 is effective in other *Lotus* species of agronomical interest (Gonnet and Díaz, 2000), we also estimated the potential benefits of hydrogen recycling carried out by *M. loti* U261H4 when in symbiosis with other 3 hosts of the same genus, namely *L. tenuis, L. subbiflorus*, and *L. uliginosus*. Similarly to what was observed in *L. corniculatus*, nodules induced by *M. loti* U261H4 engineered strain showed undetectable levels of hydrogen evolution in symbiosis with other *Lotus* species.
these 3 hosts, whereas nodules induced by the wild type strain MI U261 evolved large amounts of hydrogen in all cases (data not shown). These results indicate that *R. leguminosarum* *hup* system is expressed under symbiotic conditions in all host partners at levels high enough as to fully recycle hydrogen evolved by nitrogenase. Despite full hydrogen recycling, there were no significant differences between the wild type and U261H4 engineered strain neither in shoot dry weight nor in N content in symbiosis with *L. tenuis*, *L. subbiflorus*, *L. uliginosus* at 60 days of cultivation (Table 4), indicating a relevant role for the host on the effect of hydrogen-recycling on nitrogen fixation.

**DISCUSSION**

Hydrogenase Activity and Forage Legume Productivity

Forage legumes constitute a relevant input of protein for the sustainable feeding of livestock, a crucial issue in a context of increased demand of livestock products (FAO, 2011; Michalk et al., 2019). The productivity of pastures is dependent on an adequate supply of nitrogen (Phelan et al., 2015), therefore ensuring optimal functioning of symbiotic nitrogen fixation is a relevant aspect for pasture management. This optimization includes the use of highly efficient rhizobial strains. Even the most efficient rhizobial strains still dedicate a minimum of 25% of electrons going through nitrogenase to reduce protons to hydrogen. In this work we have generated rhizobial strains effective in two forage legumes (*Lotus* and *Vicia*) and able to fully recycle the hydrogen evolved from nitrogenase activity in the nodules. Engineered strains were associated to levels of nitrogen fixation higher than the respective wild-type strains under controlled conditions. It has to be noted, however, that for *L. tenuis* and *L. subbiflorus* the data showed a higher level of variability which, along with the limited number of replicates, did not allow to ascribe statistical significance to the observed variation of the mean values.

Previous evidence had shown that *R. leguminosarum* hydrogenase system is expressed in nodules when the hydrogenase gene cluster is introduced into strains effective on *Vicia* and *Lotus* (Monza et al., 1997; Brito et al., 2000). In contrast, introduction of the same system into *Sinorhizobium meliloti* strains resulted in only residual levels of hydrogenase activity in alfalfa nodules. It was concluded that several factors, including nickel provision and effectiveness of heterologous regulators, limit the expression in different backgrounds (Brito et al., 2000). The original transposon contains a spectinomycin resistance cassette to help on the selection of transposants (Bascones et al., 2000). We have also generated derivative strains in which such marker has been deleted, thus resulting in engineered strains in which only rhizobial DNA was present. As expected, removal of SpcR tag did not change the symbiotic performance of the strains. These strains should be more amenable to its use as inoculants.

The results presented in this work indicate that the introduced ability of hydrogen recycling has the potential to increase nitrogen fixation in forage legumes, confirming the notion of hydrogen evolution as a source of inefficiency of nitrogen fixation by the *Rhizobium*-legume symbiosis (Evans et al., 1988). There was scarce previous information on the effect of hydrogen recycling on forage legumes. The presence of hydrogenase genes in rhizobia is highly variable depending of the system (Ruiz-Argüeso et al., 2000), and in fact there are no reports on the presence of functional *hup* gene clusters in rhizobia isolated from forage legumes such as clover or alfalfa. Even in *R. leguminosarum* by viciea the presence of hydrogenase system is highly uncommon (Fernandez et al., 2005), and so the actual evolutionary advantage of possessing such system is a matter of controversy. Now we demonstrate for the first time that the advantages of hydrogen recycling can be exploited also in forage legumes, at least under controlled conditions. The results

### TABLE 3 | Effect of the introduction of *hup* genes on symbiotic performance of *M. loti* U261 in association with *L. corniculatus* plants.

| Strain        | 40 days | 60 days | 70 days |
|---------------|---------|---------|---------|
|               | N content | DW    | N content | DW    | N content | DW    |
| U261          | 114 ± 15a | 32 ± 8a | 552 ± 25a | 138 ± 33a | 253 ± 22a | 100 ± 24a |
| U261H4        | 148 ± 20a | 41 ± 10a | 627 ± 32b | 128 ± 31a | 527 ± 26b | 161 ± 39a |

*U261 strain in association with the indicated host legume.*

### TABLE 4 | Effect of the introduction of *hup* genes on symbiotic performance of *M. loti* U261 strain in association with the indicated host legumes.

| Strain     | 40 days | 60 days | 70 days |
|------------|---------|---------|---------|
|            | N content | DW    | N content | DW    | N content | DW    |
| U261       | 604 ± 180a | 155 ± 96a | 533 ± 160a | 116 ± 59a | 504 ± 45a | 126 ± 26a |
| U261H4     | 864 ± 210a | 201 ± 56a | 766 ± 180a | 187 ± 21a | 457 ± 60a | 127 ± 19a |

*Values are the mean of 3 replicates ± S.D. Values following by different letters indicate significant differences (p < 0.05) according to a Fisher's L.S.D. test.*
obtained also indicate that both the host species and the rhizobial strain condition the existence of significant increases of fixed nitrogen associated to hydrogenase activity. Factors proposed to account for the beneficial effect of hydrogen recycling are the protection of nitrogenase against oxygen and against the detrimental effect of hydrogen, and the availability of additional energy for nitrogen fixation (Palacios et al., 2005). Additional work is required to elucidate the reasons for the observed effect of both the rhizobial background and host legume. It has to be noted that nickel provision and effectiveness of heterologous regulators for expression of hydrogenase genes have been cited before as potential limiting factors (Brito et al., 2000).

**Hydrogen and Soil Fertility**

The role of hydrogen evolution and hydrogen recycling in plant productivity is not a simple question. Results presented here refer to the levels of nitrogen fixed by the inoculated legume, and have been obtained in an artificial substrate without the soil microbiome. From a more integrated point of view, the positive role of hydrogen recycling in the productivity of legumes must be considered along with the potential effect of hydrogen evolved from nodules in the fertility of surrounding soil. It has been shown that the presence of hydrogen evolving nodules in legume plants inoculated with hydrogenase-negative rhizobia results in an increase of the ratio of H2-oxidizing bacteria in soil surrounding nodules, as could be expected from the presence of an additional energy substrate in the environment. In fact, a "hydrogen fertilization effect" has been proposed as a factor contributing to the observed benefit of legumes on the following crop in rotation practices (Golding and Dong, 2010). The direct addition of hydrogen to soils resulted in increases in tiller/plant and plant dry weight in barley (Dong et al., 2003). There is also one report indicating that barley crop following soybean inoculated with Hup−, hydrogen evolving *B. japonicum* strain resulted in higher grain yield than in the case of isogenic, hydrogen recycling strain (Dean et al., 2006). In that work an increase of hydrogen oxidation activity in soils surrounding Hup− nodules was also detected. Changes in soil microbiome around nodules producing hydrogen are expected, and it has been recently shown that exposure of soil to hydrogen levels similar to those on the soil-nodule interface indeed causes changes in soil microbiome structure and function, including the stimulation of carbon turnover (Khdhiri et al., 2017). All these data indicate that the potential role of hydrogen metabolism in plant productivity should be analyzed from an integrated perspective. The hydrogen-recycling strains effective in forage legumes developed in this work can be used to further study the effect of hydrogen metabolism on mixed grass-legume pastures in which the potential role of hydrogen in stimulating other crops can be analyzed in the same season as legume symbiosis occurs, so a more integrated study of the effect of nitrogen fixation on mixed pasture production can be carried out.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

**AUTHOR CONTRIBUTIONS**

MS, AU, and SM performed experiments and analyzed data. JM and JS designed experiments, validated data and review the manuscript. JP designed experiments, validated data, wrote draft and reviewed the manuscript. All authors contributed to the article and approved the submitted version.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fagro.2021.661534/full#supplementary-material

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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