

Carbon Requirements of Leguminous Plants Receiving Different Nitrogen Nutrition

W. MERBACH

Research Centre for Soil Fertility of the Academy of Agricultural Sciences,
Müncheberg /GDR/

There are different opinions concerning the carbon requirements of legumes at symbiotic dinitrogen fixation in comparison to mineral nutrition. Besides information on equal C-demands /HARDY and HAVELKA, 1975/ results were published about higher energy requirements of air dinitrogen binding by legume-Rhizobium symbioses /SILSBURY, 1977/. In the present paper the carbon consumption of N_2 fixation and nitrate assimilation is compared taking white lupin /Lupinus albus L./ as an example in experiments carried out at the University of Halle.

Materials and methods

White lupins /var. kievskij mutant/ were cultivated in Mitscherlich vessels containing soil as substrate until they had 8-10 leaves each /MERBACH, 1982/. After this the plants were kept for several days under the conditions of 12 h light /40 klx/ and 12 h dark in order to obtain a steady state balance. According to MCCREE and SILSBURY /1978/ in this case the following equation is valid:

$$D_O = \Delta TS + N_W + N_E$$

where: D_O = apparent CO_2 assimilation in light /mg CO_2 influx per light period of 12 h/;

ΔTS = dry matter increase per day in CO_2 equivalents;

N_E = maintenance respiration /mg CO_2 efflux per dark period of 12 h after prolonged darkness of > 24 h and consequently after consumption of short time assimilate reserves/ and

N_W = growth respiration /use for biosynthesis processes/.

From these components D_O , N_E and ΔTS were determined experimentally, N_W was calculated after $N_W = D_O - (N_E - \Delta TS)$. The estimation of D_O and N_E was carried out by means of CO_2 gas exchange analysis /IRGA method, open system/. Root respiration / N_{WR} /, too, was measured. ΔTS was obtained as the difference between the plant dry matter yield at the termination and beginning of the experiment /preharvest as starting point/. In order to convert

ΔTS into CO_2 equivalents the carbon content of dry matter was determined using the method described by JACKSON /1958/. The carbon content of the lupin dry matter amounted to 39.1% and consequently, the conversion factor from dry matter to CO_2 was 1.43.

Results

As it is evident from Table 1, the N_2 fixing plants showed a significantly higher apparent CO_2 assimilation $/D_O/$, growth respiration $/N_W/$, and

Table 1

Apparent CO_2 assimilation $/D_O/$, maintenance respiration $/N_E/$, growth respiration $/N_W/$, root respiration $/N_{Wu}/$, and C-demands of the N-assimilation at different nitrogen nutrition /mean values from 6 replications/

Components	NO_3^- nutrition /0.4 g N as $NaNO_3$ per pot/	N_2 fixation without N, inoculation with Rhizobium lupini	$/N_2-NO_3^-/$
D_O mg $CO_2/12h/^{-1}.g^{-1}\Delta TS$	2135 /100/	2365 /111/ ^x	230
N_E mg $CO_2/12h/^{-1}.g^{-1}\Delta TS$	230 /100/	232 /101/	2
N_W mg $CO_2/12h/^{-1}.g^{-1}\Delta TS$	312 /100/	527 /169/ ^x	215
N_{Wu} mg $CO_2/12h/^{-1}.g^{-1}\Delta TS$	236 /100/	461 /195/ ^x	225
g C per g N	3.6 /100/	7.0 /194/ ^x	3.4

x = vs. NO_3^- variant with $\alpha = 0.05$ significant /variance analysis, t-test, single calculation/

root respiration $/N_{Wu}/$ than the nitrate assimilating plants while synthesizing 1 g dry matter. In contrast, the values of maintenance respiration $/N_E/$ did not differ between the two variants. It is interesting to note that the increases $/N_2 - NO_3^-/$ in D_O , N_W and N_{Wu} /Table 1/ were nearly equal.

The C demand per g of fixed N_2 /7 g/ was much higher than the C requirement per g of assimilated nitrate-N /3.6 g/. Similar information is given by other investigators /PATE et al., 1979; MINCHIN et al., 1980/. Consequently, the actually assayed carbon requirement for symbiotic N_2 fixation was higher than the theoretic values calculated to 2.57 until 4 g C /PHILLIPS, 1980; HARDY and HAVELKA, 1975; SHANMUGAN et al., 1978/. This is caused mainly by the energy demands, necessary for nodule formation and maintenance /MERBACH, 1982/. The dinitrogen fixing plant needs consequently more energy /carbon/ than the nitrate assimilating plant to synthesize the same amount of substance although NO_3^- nutrition demands relatively much energy. Therefore effective N_2 fixation requires essentially sufficient assimilate providing. This fact has to be considered at the development and selection of more productive Rhizobium-plant-symbioses.

References

- HARDY, R. W. F. and HAVELKA, U. D., 1975. Nitrogen fixation research: a key to world food. *Science*. 188. 633-643.
- JACKSON, M. L., 1958. Soil chemical analysis. Englewood Cliffs. 211-214.
- MCCREE, K. J. and SILSBURY, J. H., 1978. Growth and maintenance requirement of subterranean clover. *Crop Sci.* 18. 13-18.
- MERBACH, W., 1982. Untersuchungen über Stickstoffumsatz und symbiontische N₂-Fixierung bei Körnerleguminosen. Dissertation B Univ. Halle.
- MINCHIN, F. R., SUMMERFIELD, R. J. and NEVES, M. C. P., 1930. Carbon metabolism, nitrogen assimilation and seed yield of cowpea /*Vigna unguiculata* L. Walp./ grown in an adverse temperature regime. *J. exp. Bot.* 31. 1327-1345.
- PATE, J. S., LAYZELL, D. B. and ATKINS, C. A., 1979. Economy of carbon and nitrogen in nodulated and non nodulated /NO₃ grown/ legume. *Plant Physiol.* 64. 1083-1083.
- PHILLIPS, D. A., 1980. Efficiency of symbiotic fixation in legumes. *Annu. Rev. Plant Physiol.* 31. 29-44.
- SHANMUGAN, K. T. et al., 1978. Biological nitrogen fixation. *Annu. Rev. Plant Physiol.* 29. 263-276.
- SILSBURY, J. H., 1977. Energy requirement for symbiotic nitrogen fixation. *Nature*. 267. 149-150.