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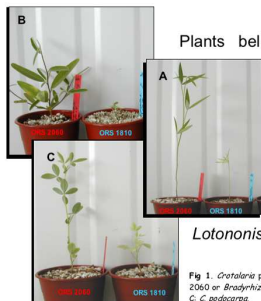
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The *Methylobacterium nodulans* / *Crotalaria podocarpa* symbiosis: a classic process for an original model

A. Renier^{1*}, S. Rapior²

¹Laboratoire des Symbioses Tropicales et Méditerranéennes (LSTM), UMR 113, Campus International de Baillarguet, 34398 Montpellier cedex 5, France.
²Laboratoire de Botanique, Phytochimie et Mycologie, UMR 5175 CEFE, Faculté de Pharmacie, 34093 Montpellier cedex 5, France.
*adeline.renier@mpl.ird.fr



Plants belonging to the *Crotalaria* genus are tropical leguminous mainly making symbiosis with *Bradyrhizobium*. Nevertheless, some *Methylobacterium* spp. strains were isolated from root nodules of *Crotalaria* as well as *Lotononis* (*Fabaceae*, *Crotalariae*) (1).

Fig 1. *Crotalaria* plants inoculated with either *Methylobacterium nodulans* ORS 2060 or *Bradyrhizobium* sp. ORS 1810. A: *Crotalaria glaucoidea*; B: *C. perrottetii*; C: *C. podocarpa*.

Methylobacterium strains were only isolated from three *Crotalaria* species, i.e., *C. glaucoidea* (Fig 1A), *C. perrottetii* (Fig 1B) and *C. podocarpa* (Fig 1C), and were described as a single novel species: *Methylobacterium nodulans*. The main feature of this original bacterial symbiotic partner is its ability to oxidize methanol, a methylotrophic property based on the presence of methanol dehydrogenase. During the symbiosis between *M. nodulans* and *C. podocarpa*, it has been shown that the bacterial methylotrophic property plays a major role in the symbiotic process (2).

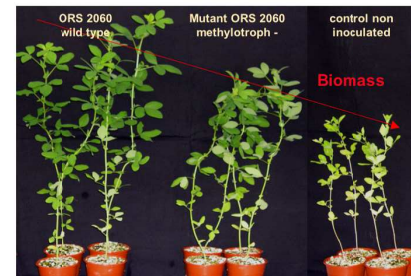
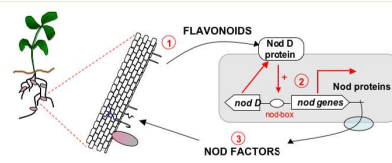


Fig 2. Comparison of growth (5 weeks after inoculation with *M. nodulans*) of *Crotalaria podocarpa*.

Nothing is known on the molecular dialogue occurring between *M. nodulans* and *C. podocarpa*. Commonly, the symbiosis between rhizobial soil bacteria and legumes was described as a multi-step process mediated by signal molecules produced from both two partners: exudation of phenolic compounds able to induce the transcription of bacterial *nod* genes leading to the biosynthesis of a bacterial signal, the nodulation factors (3).



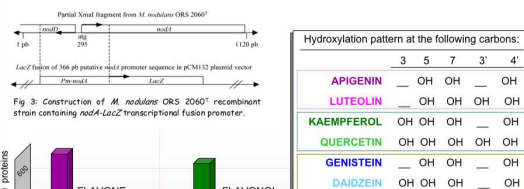
Recently Giraud et al. (3) described photosynthetic *Bradyrhizobium* ORS 278 able to nodulate *Aeschynomene indica* L. (*Fabaceae*) in a Nod factor-independent pathway, opening the possibility that rhizobia could use alternative signals to nodulate legumes.

Which signal molecules are involved in the symbiosis between *Methylobacterium nodulans* and *Crotalaria podocarpa* ?

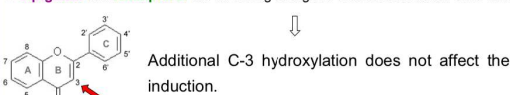
efficient flavonoid inducer of the *nod* gene expression → IDENTIFY → structure of the Nod factor produced by *M. nodulans* ORS 2060
nodulation genes operon

INDUCTION OF NOD GENES

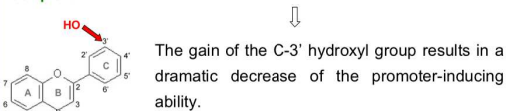
To monitor the nodulation genes expression, six flavonoids were investigated for their ability to induce the *nodA* promoter of the recombinant strain *M. nodulans* ORS2060 *nodA-LacZ*.



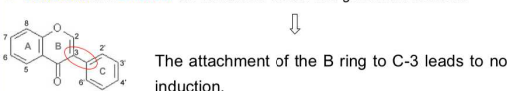
◆ Apigenin and kaempferol act as strong *nod* gene inducers on strain ORS 2060



◆ For luteolin, a 3 times decrease of activity is observed when compared to apigenin. More drastically, quercetin has no induction effect when compared to kaempferol.



◆ Genistein and daidzein are not able to induce *nod* gene in *M. nodulans*



NODULATION GENES OPERON

A 7,13-kb region in the inserted DNA fragment, showing a positive hybridization signal to the *nodA* probe from *M. nodulans* ORS 2060 was fully sequenced (Fig 4). The analysis of the nucleotide sequence revealed the existence of six entire open reading frames (ORFs).

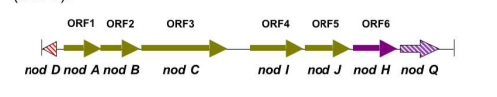


Fig 4. Organization nodulation genes in *Methylobacterium nodulans* ORS 2060 into the 7,13-kb XbaI DNA fragment. Common nodulation genes are shown in brown, specific nod genes in violet, and the nod D regulator gene in red. Partially sequence of others nod genes are indicated with hatchings.

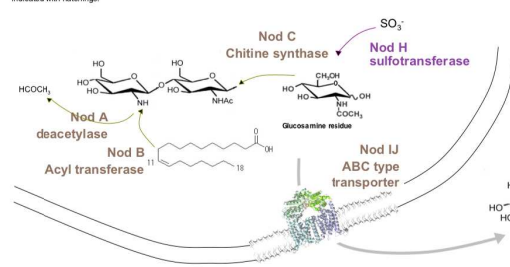


Fig 5. Functions of the different Nod proteins encoded by the genes belonging to the XbaI fragment.

NODULATION FACTORS

In order to prepare Nod factors, *M. nodulans* ORS 2060 was grown in the presence of apigenin. After induction, Nod factors were extracted from the culture supernatant and purified. HPLC analyses revealed the presence of only two fractions (Fig 6), then analysed by LC/MS.

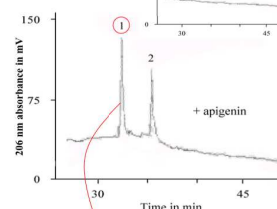


Fig 6. HPLC profile of a Nod factor extract from a apigenin-inducing *M. nodulans* ORS 2060 culture supernatant.

The positive ion spectra of compounds identified in fraction 1 show two DP5 glucosamine sulfated forms as C_{18:1} (vaccenic acid) and C_{16:0} (palmitic acid) in proportion 10/1 (Fig 7). No Nod factor was found in fraction 2.

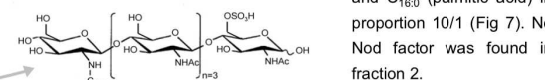


Fig 7. Structure of the major lipo-chito-oligosaccharide identified produced by *M. nodulans* ORS 2060.

◆ *Methylobacterium nodulans* is able to distinguish between flavonoids (flavone and flavonol) and isoflavonoids (isoflavone and isoflavonol). Thus, the attachment of the B ring to C-2 is of crucial importance for induction.

Commonly apigenin and surprisingly kaempferol, usually found as suppressor, are highlighted as to be the most powerful flavonoid inducers of the *nodA* promoter of *M. nodulans*.

◆ The presence of *nod* genes is checked and a DNA fragment containing 8 nodulation genes *nod DABCIJHQ* have been identified.

◆ *M. nodulans* produces one major Nod factor structure identified as to be Nod Mn-V(C_{18:1}S), suggesting a classic symbiosis model for *M. nodulans* / *C. podocarpa* association.

(1) Sy A, Giraud E, Jourand P, et al. (2001) Methylobacterium bacteria nodulate and fix nitrogen in symbiosis with legumes. *Journal of Bacteriology* 183, 214-220.

(2) Jourand P, Renier A, Rapior S, et al. (2005) Role of methylotrophy during symbiosis between *Methylobacterium nodulans* and *Crotalaria podocarpa*. *Molecular Plant-Microbe Interactions* 18, 1061-1068.

(3) Giraud E, Moulin L, Vallenet D, et al. (2007) Legumes symbioses: absence of *nod* genes in photosynthetic bradyrhizobia. *Science* 316, 1307-1312.