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Effects of long-term fertilizer applications on decomposition of $^{14}\rm C$ labelled Lolium shoots (*Lolium perenne* L.) and $^{14}\rm C$ labelled glucose

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The effects of fertilizer applications on decomposition of 14 C labelled Lolium shoots (*Lolium perenne* L.) and 14 C labelled glucose were determined in a long-term field trial included control, organic-mineral fertilizers with herbicides, organic-mineral fertilizers without herbicides, mineral fertilizers only. The incorporation of 14 C-labelled substrates to soils allows for differentiation between substrates-derived carbon and native soil organic carbon, and provides precise estimates of decomposition rates. Two types of 14 C-labelled substrates: glucose and plant residues of *Lolium perenne L*. were incubated in a silt loam Podzoluvisol to test the importance of substrate-soil relationships by decomposition of soil organic matter and turnover of C through microbial biomass.

We sampled Podzoluvisol of a long term field trial located in Moscow region, Russia. In the field trial four fertilizer systems were applied during 38 years. The first treatment (1) was control without any fertilizer; the second one included (2) organic-mineral fertilizers with herbicides; the third one (3) - organic-mineral fertilizers without herbicides; the fourth one (4) – mineral fertilizer only. The incubation was conducted under controlled conditions in closed vessels for 90 days at 20 °C. Fifteen grams of air-dried soils were weight into 250 ml glass vessels. For each of the four field treatments the following additions were tested: (i) glucose: 14,4 mg C of 14 C-labelled glucose (spe-

cific ¹⁴C activity 18,7 kBq mg^{-1C}); (ii) plant residue: 14,4 mg C of ¹⁴C-labelled plant residues (*Lolium perenne L.*) ground with ball mill (specific ¹⁴C activity 4,8 kBq mg^{-1C}); and (iii) control without glucose or plant residues. The decomposition of ¹⁴C labelled plant residue and ¹⁴C labelled glucose was estimated by measuring the ¹⁴CO₂ emissions. Small vials with 3 ml of 1.0 M NaOH were placed into the vessels for trapping CO₂. The traps were changed after 1, 2, 4, 7, 11, 19, 29, 41, 71 è 90 days (10 samplings). Biomass C in soil was analysed after 8, 19, 41 and 90 days by the fumigation-extraction method (Vance et al., 1987). Deionized water was added to the vessels as needed to keep soil moisture at 70% of WHC.

To compare the CO₂ efflux rates from field treatments having initially different C_{org} content, all results are presented as percentage of SOC. At the first day all soil treatments showed initially CO₂ efflux rate accounting for 0.15-0.18% and 0.10-0.11% of SOC day⁻¹ for glucose and plant residues, respectively. Then the CO₂evolution from soil dropped to 0.06 and 0.07% of SOC day⁻¹ for glucose and plant residues by day 3. Thereafter, the CO₂ evolution rate became lower and generally constant, remained at 0.01% of SOC day⁻¹ for both substance additions until the end of 90 day incubation. The cumulative CO₂ production over 90 days fitted very well to the double exponential model (R² >0.99 for the four soils) (Bonde et al., 1988).

The effects of four fertilizer systems on the mineralization of 14 C-labelled glucose and plant residues showed that mineralization of glucose and plant residues began immediately with the maximum rate occurring within 2 days, ranging 22.4 % and 7.89 % of added ¹⁴C activity per day for glucose and plant residues, respectively. At the end of the incubation, the cumulative ¹⁴CO₂ evolved amounted to 77 and 66 % of added C of glucose and plant residues, respectively. The decomposition rate (k) of labile ¹⁴C-labelled glucose was higher in control treatment (1) and in soil treatment with organic-mineral fertilizers with herbicides (3) (0.31, 0.29 d⁻¹, respectively) comparing with organic-mineral fertilizers with herbicides (2) and mineral fertilizer only (1) (0.24, 0.25 d⁻¹, respectively). k of labile ¹⁴C-labelled plant residues had the same tendency. In the same treatments (1,3) with addition of glucose microbial biomass C was higher during 20 days of incubation (1 mg C/1g soil for 1 and 3 treatments and less 0.8 mg C/1g soil for 2 and 4 treatments).

We determined no significant effect of organic-mineral and mineral fertilizers systems (2,3,4) on CO₂ native carbon mineralization. There was significant effect only in control comparing with fertilizer application. Initially glucose decomposition was 2.7 times faster than plant residues decomposition. The decomposition rates of labile ¹⁴C-labelled plant residues and ¹⁴C-labelled glucose were higher in control treatment and in soil treatment with organic-mineral fertilizers without herbicides and related to microbial biomass C.