

A Norleucine-resistant Mutant of *Salmonella typhimurium* with a Possible Defect in Valine Uptake or Regulation

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INTRODUCTION

All the *Salmonella typhimurium* mutants resistant to the methionine analogue norleucine described so far are also resistant to the analogues ethionine and α -methylmethionine. They can be classed either as *metA*(fbr) mutants with a homoserine-*O*-transsuccinylase, the first enzyme specific to methionine biosynthesis, resistant to feedback inhibition by methionine or α -methylmethionine, or as *metK* mutants with defects in *S*-adenosylmethionine (SAM) synthetase leading to abnormalities in methionine repression control (Lawrence, Smith & Rowbury, 1968; Chater & Rowbury, 1970; Hobson & Smith, 1973). A recently isolated norleucine-resistant mutant of a cysteine and thymine auxotroph, *cysJ538 thyA*, was found to be sensitive to ethionine. This mutant, provisionally designated *cysJ538 thyA* Nol-r, was studied in the hope that it might be a new type of methionine regulatory mutant.

METHODS

Organisms. All bacterial strains were derivatives of *Salmonella typhimurium* LT-2. *CysJ538 thyA* Nol-r, a spontaneous norleucine-resistant mutant of *cysJ538 thyA*, was provided by Dr M. C. Jones-Mortimer, Department of Biochemistry, University of Leicester, as were strains carrying *Escherichia coli* F'-factors. Hfr strains were obtained from Dr K. E. Sanderson, Department of Biology, University of Calgary, Alberta, Canada. Phage L4, a non-lysogenizing variant of P22 (Smith & Levine, 1967), was routinely used in transduction. Maintenance of stock cultures, growth of bacteria for genetical experiments, and the propagation, assay and maintenance of transducing phage were as described by Smith (1961).

Media. Oxoid nutrient agar (NA) and broth (NB) were used. The minimal medium (MM) and minimal agar (MA) were those of Smith (1961) and were supplemented as required with the following ($\mu\text{g/ml}$): DL-ethionine, 1000; α -methyl-DL-methionine, 1000; DL-norleucine, 1000; thymine, 60; L-valine, 40; other amino acids, 20.

Genetical techniques. (i) Transduction. The technique of Smith (1961) was used. (ii) Conjugation. Overnight NB cultures of recipient and donor strains were mixed in the ratio 1:1 and the mixtures incubated without shaking for 30 min. Dilutions (10^{-2}) were then made in NB with gentle mixing and 0.2 ml portions of these spread on to appropriately supplemented MA plates. Identical plates were spread with recipient and donor strains separately as controls. All plates were incubated for 40 to 48 h. (iii) Episome transfer. Both transfer and curing of episomes were as described by Hobson & Smith (1973).

Enzyme assays. Bacteria were cultured, extracts prepared and SAM synthetase activity assayed as described by Hobson & Smith (1973). The N^5N^{10} -methylene tetrahydrofolate reductase assay was that used by Whitehouse & Smith (1973).

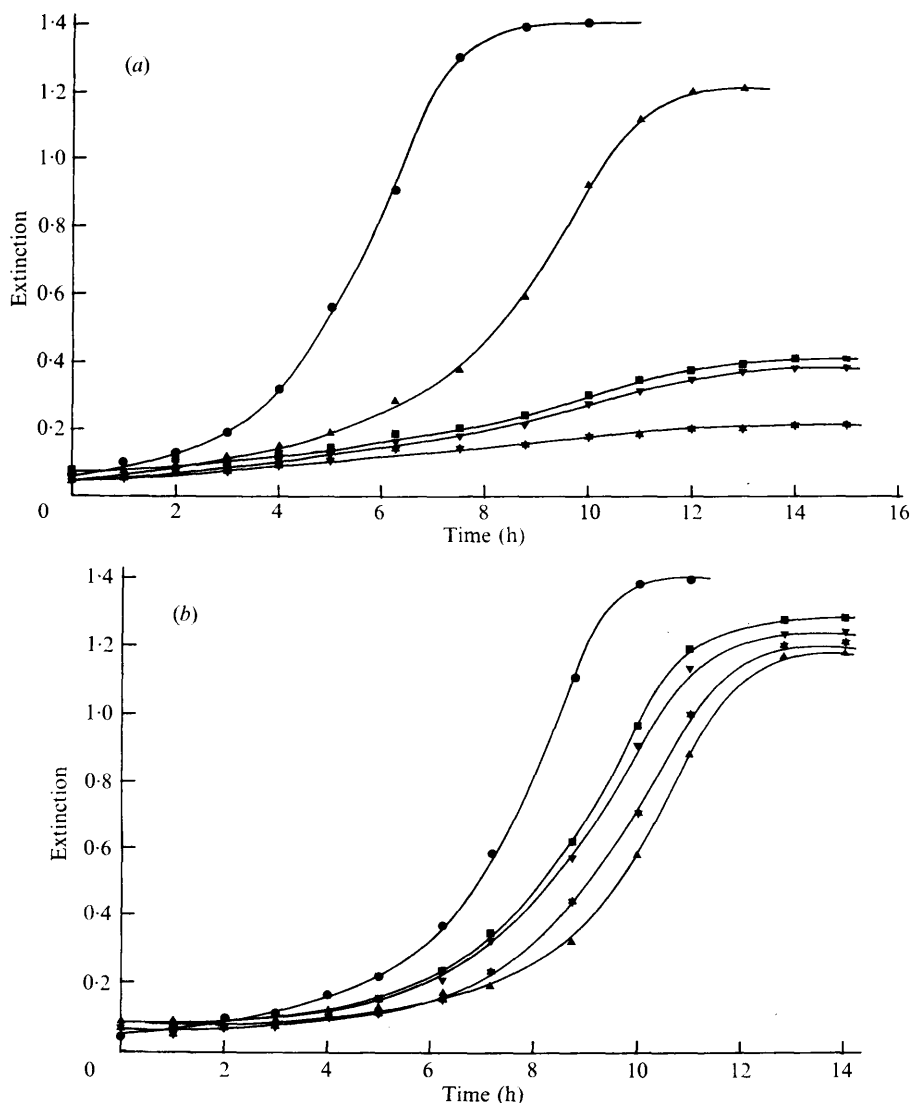


Fig. 1. The effect of methionine, valine and leucine on growth in MM + norleucine of (a) wild type; (b) *metK⁷²⁵*; (c) *cysJ538 thyA Nal-r*. All supplements were added to MM at time 0 and were: ●, no supplement; *, norleucine; ▲, norleucine + methionine; ▼, norleucine + valine; ■, norleucine + leucine. In addition, cysteine and thymine were added to *cysJ538 thyA Nal-r* cultures.

Growth experiments. Bacterial cultures were grown to stationary phase in appropriately supplemented MM, concentrated tenfold and diluted into 15 ml fresh MM supplemented as required to give an extinction of 0.05 to 0.1 measured with a Unicam SP500 spectrophotometer (650 nm, 1 cm light path). These cultures were incubated in 25 ml conical flasks on a reciprocating shaker at 37 °C. Samples (1.5 ml) were removed at approximately hourly intervals, their extinction measured as before and the samples discarded.

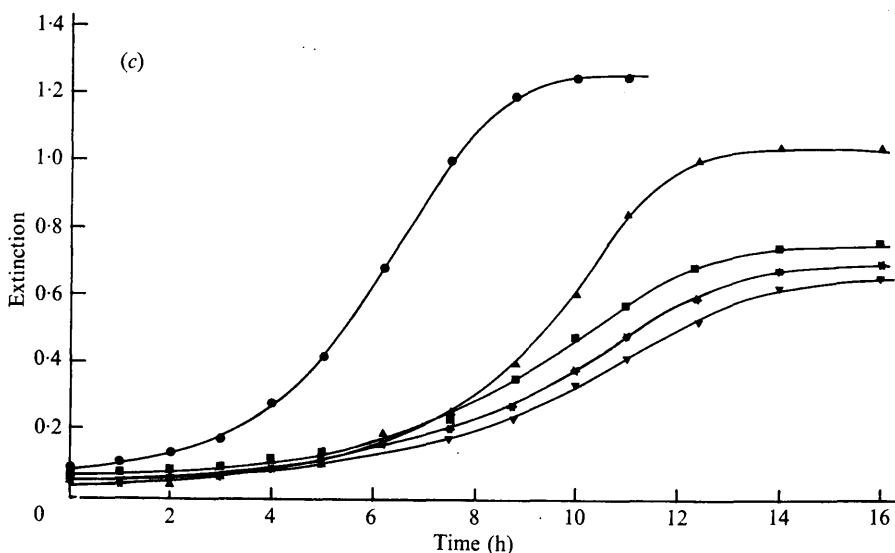


Fig. 1(c). For legend see opposite.

RESULTS AND DISCUSSION

The norleucine resistance and ethionine sensitivity of *cysJ538 thyA* Nol-r was first confirmed by growth experiments in liquid medium. The mutant was similarly shown to be sensitive to α -methylmethionine so that the Nol-r mutation is unlikely to be of the *metA*(fbr) type or to affect the methionine-specific permease because mutants of the permease are resistant to this analogue (Ayling & Bridgeland, 1972). It was possible that the mutation was in either the *metJ* or *metK* gene, both of which are involved in repression by methionine of its own synthesis, but gave a different analogue resistance pattern from previously studied mutations in these genes. *MetJ* is about 95 and 45% cotransducible with the *metB* and *metF* structural genes respectively, and *metK* is about 1% cotransducible with *serA* (Lawrence *et al.* 1968). However, in transduction crosses using *cysJ538 thyA* Nol-r as donor no linkage of Nol-r with any of these auxotrophic markers could be detected out of at least 300 transductants tested in crosses with *metB* and *metF* as recipients and at least 1000 transductants with *serA* as recipient, showing that Nol-r is not in either the *metJ* or *metK* gene. *CysJ538 thyA* Nol-r had the same SAM synthetase activity as wild-type and so cannot be defective in some subunit or cofactor of the *metK* enzyme.

In seeking a defect in methionine regulation, *cysJ538 thyA* Nol-r was shown not to excrete methionine in cross-feeding tests with a methionine auxotroph on supplemented MA (as in Lawrence *et al.* 1968) but other mutants do not excrete methionine and yet have elevated levels of methionine biosynthetic enzymes and methionine pools (Hobson, 1973). However, the *cysJ538 thyA* Nol-r *N⁵N¹⁰*-methylenetrahydrofolate reductase (involved in the terminal step of methionine biosynthesis) was found to be neither derepressed nor resistant to repression by 10 mM-L-methionine, and its methionine pool was the same as that of wild type as determined by a method involving a bioassay of methionine in cell extracts (Hobson, 1973). Thus this Nol-r mutation, unlike all others which lead to norleucine resistance, does not affect methionine regulation.

Another explanation for the Nol-r phenotype could be interference with uptake of norleucine. This analogue is transported at low efficiency by the methionine permease (Ayling &

Bridgeland, 1972) and norleucine, leucine and isoleucine compete with valine for uptake by *Escherichia coli*, suggesting common uptake systems for these compounds (Cohen & Rickenberg, 1956). In addition, if part of the inhibition of wild type by norleucine results from interference with synthesis or utilization of another amino acid besides methionine, the Nol-r mutation could cause overproduction of the relevant amino acid. For example, Trupin, Dickerman, Nirenberg & Weissbach (1966) showed that norleucine could be activated by tRNA^{Leu}.

To test these hypotheses, the reversal of norleucine inhibition of wild-type growth by growth concentrations of methionine, valine and leucine was tested in MM. All three amino acids stimulated growth, methionine having the largest effect, their combined effects almost completely reversing the inhibition (Fig. 1a). Thus norleucine must interfere with either uptake, synthesis or utilization of all three compounds. As expected, methionine did not stimulate growth of *metK*⁷²⁵ (a mutant which overproduces methionine) in norleucine, but valine and leucine increased growth as with wild type (Fig. 1b). However, when the growth experiments were repeated with *cysJ538 thyA* Nol-r, valine had no stimulatory effect on growth in norleucine while methionine and leucine retained their stimulation (Fig. 1c). Thus in this mutant the interference of norleucine with valine metabolism seems to be blocked. This could mean that *cysJ538 thyA* Nol-r lacks a valine uptake system, overproduces valine, or somehow affects the utilization of this amino acid, perhaps by altering the tRNA synthetase.

Knowledge of the genetic map location of the Nol-r mutation could help with interpretation of the nature of the defect it causes, e.g. if it mapped in the *ilv* (isoleucine/valine) cluster. Preliminary conjugation crosses were carried out using *cysJ538 thyA* Nol-r as a recipient and HfrB2*leu* and HfrK3*his* as donors with separate selection for *cys*⁺ and *thy*⁺. *CysJ* and *thyA* map at about 90 and 92 min respectively (Sanderson, 1972). With HfrB2*leu*, whose origin is at about 55 min, 95 to 100% linkage of Nol-r to both markers was obtained. The HfrK3*his* origin is at about 93 min and transfer occurs with *thyA* as a very late marker. With this Hfr strain few *thy*⁺ recombinants were obtained, but 30% linkage with Nol-r was found whereas linkage between *cysJ* and Nol-r was low (7%). These results suggest that Nol-r lies between *cysJ* and the HfrK3 origin. This is in agreement with results obtained by transferring F'-factors to *cysJ538 thyA* Nol-r. Two episomes were used: KLF16 which carries the region of the genome from *metC* at about 102 min to just beyond *thyA* but not extending into the *cys* cluster, and another F'-factor carrying the region from about 94 to 90 min, that is including the *cys* genes. Each episome was transferred with selection for *thy*⁺ and in each case the heterogenotes were found to be sensitive to norleucine. Curing of the episome resulted in the simultaneous appearance of norleucine resistance and a thymine requirement. Thus Nol-r is recessive and must map in the region covered by both episomes between the *cys* cluster and *lysA* (94 min). However, when phage grown on *cysJ538 thyA* Nol-r, or the *cys*⁺ or *cys*⁺*thy*⁺ derivatives, was used to transduce to prototrophy mutants in the *cys* cluster and also in *thyA*, *argB* and *lysA* which are all cotransduced by phage P22, no norleucine-resistant transductants were obtained from at least 300 scored in each cross. Thus the most likely map position for the Nol-r mutation is between the two linkage groups, near *recA* (Sanderson, 1972).

In *Escherichia coli*, mutants which over-produce valine map in the *ilv* cluster, in genes concerned with branched-chain amino acid synthetases or in *azl* (azaleucine resistance), all of which map away from the postulated Nol-r location (Umbarger, 1971; Taylor & Trotter, 1972). However, it is possible that the Nol-r mutation is in a different, previously unrecognized valine regulatory gene. Nol-r could possibly reduce incorporation of valine into protein, but

it maps away from the known tRNA synthetase genes and genes affecting tRNA itself. The other possibility is that *cysJ538 thyA* Nol-r is a permease mutant. Two genes involved in uptake of branched-chain amino acids have been mapped in *E. coli*: *brnP* and *brnQ* mapping at about 5 and 10 min respectively (Guardiola & Iaccarino, 1971). *CysJ538 thyA* Nol-r could be defective in another valine permease or could affect a reaction coupled with valine transport (Kaback, 1972).

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