

# Role of neuronal and inducible nitric oxide synthases in the guinea pig ileum myenteric plexus during *in vitro* ischemia and reperfusion

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## Abstract

**Background** Intestinal ischemia and reperfusion (I/R) injury leads to abnormalities in motility, namely delay of transit, caused by damage to myenteric neurons. Alterations of the nitrergic transmission may occur in these conditions. This study investigated whether an *in vitro* I/R injury may affect nitric oxide (NO) production from the myenteric plexus of the guinea pig ileum and which NO synthase (NOS) isoform is involved. **Methods** The distribution of the neuronal (n) and inducible (i) NOS was determined by immunohistochemistry during 60 min of glucose/oxygen deprivation (*in vitro* ischemia) followed by 60 min of reperfusion. The protein and mRNA levels of nNOS and iNOS were investigated by Western-immunoblotting and real time RT-PCR, respectively. NO levels were quantified as nitrite/nitrate. **Key Results** After *in vitro* I/R the proportion of nNOS-expressing neurons and protein levels remained unchanged. nNOS mRNA levels increased 60 min after inducing ischemia and in the following 5 min after reperfusion. iNOS-immunoreactive neurons, protein and mRNA levels were up-regulated during the whole I/R period. A significant increase of nitrite/nitrate levels was observed in the first 5 min after inducing I/R and was significantly reduced by N<sup>ω</sup>-propyl-L-arginine and 1400 W, selective inhibitors of nNOS and iNOS,

respectively. **Conclusions & Inferences** Our data demonstrate that both iNOS and nNOS represent sources for NO overproduction in ileal myenteric plexus during I/R, although iNOS undergoes more consistent changes suggesting a more relevant role for this isoform in the alterations occurring in myenteric neurons following I/R.

**Keywords** guinea pig ileum, *in vitro* ischemia/reperfusion, myenteric plexus, nitric oxide, nitric oxide synthase.

## INTRODUCTION

Intestinal ischemia/reperfusion (I/R) injury represents an important clinical problem which may occur as a consequence of embolism, arterial or venous thrombosis, shock,<sup>1</sup> intestinal transplantation, necrotising enterocolitis in the human premature newborn or chronic inflammatory diseases.<sup>2–4</sup> I/R injury causes severe damage particularly to the mucosal and muscle layers and neurons.<sup>5–7</sup> The functional consequences of this scenario include impairment of nutrient absorption, altered intestinal barrier function against bacterial translocation and disturbed motility.<sup>5,6,8</sup> Adaptive changes of enteric neuronal circuitries modulating the intestinal motor function have been documented after both *in vivo* and *in vitro* I/R conditions.<sup>9–11</sup> Among the major enteric neurotransmitter pathways, the nitrergic transmission has been suggested to participate in the motor disturbances caused by an I/R injury to the gut. In physiologic conditions, nitric oxide (NO) plays a pivotal role in the inhibitory regulation of peristalsis.<sup>12,13</sup> NO is synthesized by the neuronal (n), endothelial (e) and inducible (i) isoforms of nitric oxide synthase (NOS), all of which have been localized in

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myenteric neurons of different species.<sup>14–17</sup> nNOS is constitutively expressed in myenteric neurons<sup>15–17</sup> and is the predominant source for the generation of NO mediating non-adrenergic non-cholinergic inhibition of smooth muscle contraction.<sup>18</sup> In myenteric neurons, expression of iNOS, but not of nNOS or eNOS, is enhanced during disease states, such as intestinal inflammation<sup>19,20</sup> and diabetes.<sup>21</sup> In these conditions, large amounts of NO may cause damage to different cell types, including neurons, by the formation of peroxynitrite and nitrotyrosine.<sup>20,22</sup> Previous studies suggest that the activity of the diverse NOS isoforms may be differentially regulated during intestinal I/R, depending on the different experimental model and on the degree of damage. In the rat ileum, after a severe *in vivo* I/R injury, iNOS up-regulation was correlated with a decrease of the intestinal transit.<sup>23</sup> Conversely, in the same animal model, a mild I/R injury induced a reduction of nNOS activity, which was correlated with a transitory impairment of the non-adrenergic non-cholinergic inhibitory response.<sup>24</sup> Such changes of the intestinal motor function may rely upon alterations of NOS containing myenteric neurons. In the rat ileum, a significant increase in the number of myenteric neurons expressing NOS immunoreactivity has been demonstrated after an *in vivo* I/R injury.<sup>25</sup> Administration of *N* $\epsilon$ -nitro-L-arginine methyl ester, a non-selective NOS inhibitor, prevented the increase of NOS immunoreactivity induced by I/R, suggesting the occurrence of a positive feedback between NO production and NOS expression. In the guinea pig ileum, mild *in vivo* I/R damage induced swelling and distortion of dendrites of NOS-expressing descending inhibitory motor neurons.<sup>26</sup>

In the present study, to further investigate the involvement of the nitrergic transmission during I/R, we evaluated whether an *in vitro* I/R injury may acutely affect NO production from the guinea pig ileum muscle wall. In addition, the hypothesis that nNOS and iNOS isoforms may represent sources for NO in these experimental conditions was investigated. The potential different roles of the two isoforms was evaluated using molecular biology, morphological and functional approaches.

## MATERIALS AND METHODS

### Animals and tissue preparation

Male Dunkin-Hartley guinea pigs (Harlan Italy, Correzzana, Monza, Italy), weighing between 300 and 350 g, were housed in groups of four under controlled environmental conditions (temperature  $22 \pm 2$  °C; relative humidity 60–70%) with free access to a standard diet and water, and were maintained at a

regular 12/12-h light/dark cycle. Principles of good laboratory animal care were followed and animal experimentation was in compliance with specific national and international laws and regulations. Animals were sacrificed by decapitation and 8 cm segments of the ileum, approximately 5 cm oral to the ileocaecal junction, were rapidly excised and rinsed with a physiological ice-cold Tyrode's solution [composition (mmol L<sup>-1</sup>): 137 NaCl; 2.68 KCl; 1.8 CaCl<sub>2</sub>·2H<sub>2</sub>O; 2 MgCl<sub>2</sub>; 0.47 NaH<sub>2</sub>PO<sub>4</sub>; 11.9 NaHCO<sub>3</sub>; 5.6 glucose]. Whole wall intestinal segments were fixed and stored for successive immunohistochemistry experiments, as described below. Overflow experiments were carried out on mucosa-deprived ileal segments. Western immunoblot and real time RT-PCR studies were conducted using preparations consisting of external longitudinal muscle layer segments with attached myenteric plexus (LMMP) obtained immediately after excision of ileal segments.

*In vitro* ischemic conditions were reproduced by suspending intestinal preparations in 3 mL organ baths superfused with an oxygen and glucose deprived Tyrode's solution. Oxygen deprivation was obtained by bubbling the perfusing medium with a mixture of N<sub>2</sub>-CO<sub>2</sub> (95–5%). The effect of reperfusion was evaluated by substituting the hypoxic/hypoglycaemic medium with a normal oxygenated glucose-containing Tyrode's solution. During overflow studies, ileal segments were exposed to *in vitro* ischemic conditions for 60 min and subsequently reperfused for another 60 min period. For immunohistochemistry, Western immunoblot, real time RT-PCR studies, intestinal specimens after being perfused with an oxygenated Tyrode's solution maintained at 36.5 °C for a 60 min equilibration period, were exposed to *in vitro* ischemic conditions for 5, 30 or 60 min and then reperfused for 5, 30 and 60 min. This time course for glucose/oxygen deprivation was chosen given that enteric neurons are not irreversibly damaged after 60 min ischemia.<sup>8</sup> Control experiments of Western immunoblotting and real time RT-PCR were carried out by exposing intestinal specimens to a normal glucose-containing and oxygenated medium for 5, 30, 60, 65, 90, and 120 min after the equilibration period. Tissue samples were either stored at -70 °C for successive Western immunoblotting assay or stored in a preserving solution for downstream real time RT-PCR (RNA later™; Ambion Life Technologies Italia, Monza, Italy).

### Immunohistochemistry

Segments of the guinea pig ileum were fixed for 4 h at room temperature (RT) in 0.2 mol L<sup>-1</sup> sodium phosphate-buffer (PBS: 0.14 mol L<sup>-1</sup> NaCl, 0.003 mol L<sup>-1</sup> KCl, 0.015 mol L<sup>-1</sup> Na<sub>2</sub>HPO<sub>4</sub>, 0.0015 mol L<sup>-1</sup> KH<sub>2</sub>PO<sub>4</sub>) pH 7.4, 4% formaldehyde and 0.2% picric acid. Preparations were then cleared of fixative and stored at 4 °C in PBS containing 0.05% thimerosal. LMMP whole-mount ileal preparations were then prepared according to the method of<sup>27</sup> Briefly, after blocking aspecific sites with PBS, 1% Triton X-100 and 10% normal horse serum (NHS) (Euroclone, Celbio, Milan, Italy) for 1 h, preparations were incubated with optimally diluted primary antibodies (Table 1). Double labelling was performed during consecutive incubation times: firstly, the primary antibody raised against either iNOS or nNOS was added overnight at 4 °C, then incubation with secondary antibodies followed for 2 h at RT. Whole mounts were successively incubated overnight at 4 °C with a biotinylated antibody to HuC/D (neuronal cell marker) or with an antibody to S100 (glial cell marker) (Table 1). Incubation for 2 h with either streptavidin conjugated Cy3 or with an appropriate secondary antibody was then performed at RT. Double labelling with iNOS and nNOS was performed following the same procedures, by adding the two primary antibodies consecutively. Preparations were mounted

**Table 1** Primary and secondary antisera and their respective dilutions used for western blot (WB) assay and immunohistochemistry (HC)

Antiserum	Dilution (WB)	Dilution (HC)	Source	Host species
<b>Primary antisera</b>				
iNOS	1 : 200	1 : 250	Santa Cruz (sc8310; H-174)	Rabbit
iNOS	–	1 : 50	AbCam (ab49999)	Mouse
nNOS	1 : 500	1 : 250	Millipore (AB1529)	Sheep
HUC/D	–	1 : 100	Molecular Probes, Invitrogen (A-21272)	Mouse
S-100	–	1 : 1000	Dako (Z0311)	Rabbit
$\alpha$ -tubulin	1 : 1000	–	Sigma-Aldrich (T-6199 DM-1A)	Mouse
<b>Secondary antisera &amp; streptavidin complexes</b>				
Anti-sheep FITC	–	1 : 150	Millipore (AP147F)	Rabbit
Anti-rabbit Alexa Fluor 488	–	1 : 200	Molecular Probes (A21206)	Donkey
Anti-mouse Alexa Fluor 488	–	1 : 300	Molecular Probes (A21202)	Donkey
Cy3-conjugated streptavidin	–	1 : 500	Amersham (PA43001)	–
Anti-sheep IgG horseradish peroxidase-conjugated	–	1 : 20 000	Millipore (AP147F)	Rabbit
Anti-rabbit IgG horseradish peroxidase-conjugated	1 : 10 000	–	Amersham (NA934)	Donkey
Anti-mouse IgG horseradish peroxidase-conjugated	1 : 10 000	–	Millipore (AP126P)	Millipore (AP147F)

Supply companies: Amersham, GE Healthcare, Buckinghamshire, UK; Abcam, Cambridge, UK; Millipore Co, Billerica, MA, USA; Dako, DK-2600 Glostrup, Demark; Molecular Probes, Invitrogen, Carlsbad, CA, USA; Santa Cruz Biotechnology, Inc, CA, USA; Sigma-Aldrich, Milano, Italy.

onto glass slides, using a mounting medium with DAPI (Vectashield<sup>®</sup>; Vector Lab, Burlingame, CA, USA). Neuron counts were made on HuC/D stained LMMPs, obtained from at least four animals, digitized by capturing as many as 40X objective microscope fields (0.287 mm<sup>2</sup>) as possible (10–15 fields). The total value was divided by the total image field area considered and expressed as the number of cell bodies mm<sup>-2</sup>. To establish the proportion of iNOS and nNOS expressing myenteric neurons, quantitative analysis of double fluorescently labelled ileal whole mounts was performed as previously described.<sup>27,28</sup> Negative controls and interference control staining was evaluated by omitting one or both of the primary antibodies, or one of the secondary antibodies. Preparations were analyzed by confocal microscopy on a Leica TCS SP5 confocal laser scanning system (Leica Microsystems GmbH, Wetzlar, Germany) and pictures were processed using Adobe-Photoshop CS2.0 software.

### Western immunoblot analysis of iNOS and nNOS

iNOS and nNOS protein level analysis was carried out starting from LMMPs preparations with successive centrifugations according to the method described<sup>27</sup> The purified membrane fraction and the supernatant were boiled for 2 min after dilution with Laemmli sample buffer<sup>29</sup> and processed as described elsewhere.<sup>27</sup> Membranes were incubated with optimally diluted primary and horseradish peroxidase-conjugated secondary antisera (Table 1). Blots for iNOS and nNOS were developed using an enhanced chemiluminescence technique (ECL advance Amersham Pharmacia Biotech, Cologno Monzese, Italy). Signal intensity was quantified by densitometric analysis using the NIH image software 1.61 (downloadable at <http://rsb.info.nih.gov/nih-image>). In each membrane  $\alpha$ -tubulin was monitored and used as protein loading control. Experiments were performed at least four times for each different preparation. The effect of either *in vitro* I/R or of superfusion with a normal glucose-containing oxygenated medium on protein levels was expressed as the percentage variation vs values obtained in preparations exposed to normal metabolic conditions and collected at the end of the equilibration period. Specificity of iNOS and nNOS primary antibodies was evaluated by testing their selectivity in RAW 264.7 macrophage cells and in the rat hippocampus (data not shown), respectively, and by omission of the primary antibody.

### Real time quantitative RT-PCR

Total RNA was extracted from LMMPs preparations with TRIzol (Invitrogen) and treated with DNase I (DNase Free, Ambion) to remove any traces of contaminating DNA. cDNA was obtained retrotranscribing 2  $\mu$ g of total RNA using the High Capacity cDNA synthesis kit (Applied Biosystems, Life Technologies, Grand Island, NY, USA). Quantitative RT-PCR was performed on the Abi Prism 7000 real-time thermocycler (Applied Biosystems) with Power Sybr Green Universal PCR Master Mix (Applied Biosystems) following manufacturer's instructions. Primers were designed using Primer Express software (Applied Biosystems) on the basis of available sequences deposited in public database. Primer sequences were: n-NOS, 5'-ACGGAGCGTGGCTACAGTTT-3'; 5'-TCCTTGATGTCACGCACAATTT-3'; iNOS: 5'-GCAGCAGCGGCTTCACA-3'; 5'-ACATCCAAACAGGAGCGTCAT-3';  $\beta$ -actin 5'-ACGGAGCGTGGCTACAGTTT-3'; 5'-TCCTTGATGTCACGCACAATTT-3'; GAPDH, 5'-CGGATTTGGCCGTATTGG-3'; 5'-AATATCACCTTTGCCAGACATGAA-3'. For quantitative RT-PCR a final concentration of 100 nmol L<sup>-1</sup> for each primer was used. Primers were designed to have a similar amplicon size and similar amplification efficiency as required for the utilization of the  $\Delta\Delta$ Ct method to compare gene expression.<sup>30</sup>  $\beta$ -actin and GAPDH were used as housekeeping genes. Experiments were performed at least four times for each different preparation. The effect of either *in vitro* I/R or of superfusion with a normal glucose-containing oxygenated medium on mRNA levels was expressed as the percentage variation vs values obtained in preparations exposed to normal metabolic conditions.

### Overflow study

NO overflow was accomplished using mucosa-deprived ileal specimens. The submucosal layer was not routinely removed from the intestinal wall since preliminary experiments demonstrated that the release of NO oxidation products from the submucosal plexus was undetectable by our method of analysis. Silk ligatures were applied to each end of 1–2 cm long strips; one end was attached to a rigid support and the other to an isotonic transducer (load 1 g) in 3 mL organ baths perfused at a constant rate of 1 mL min<sup>-1</sup> with Tyrode's gassed with O<sub>2</sub>-CO<sub>2</sub> (95–5%) and maintained at 36.5 °C. After a 60 min equilibration period, three samples were

collected over a period of 5 min for each sample and considered as controls, subsequently *in vitro* I/R conditions were applied, as described above. Test drugs were added to the superfusion medium after collection of the third control sample until the end of the experiment. Equilibration periods of 20 min were allowed after addition of  $N^{\omega}$ -propyl-L-arginine ( $1 \mu\text{mol L}^{-1}$ ) and 1400 W ( $10 \mu\text{mol L}^{-1}$ ), selective inhibitors of nNOS and iNOS, respectively. A single drug concentration was tested on each intestinal preparation. After the equilibration period with the appropriate test drug, three samples were collected each over a period of 5 min, before inducing *in vitro* I/R. At the end of the experiment, ileal samples were blotted and weighed ( $189.73 \pm 5.34 \text{ mg}$ ;  $n = 33$ ).

Samples were assayed for NO by means of the nitrite method based on the Griess reaction.<sup>31</sup> Briefly, samples were incubated with Griess reagent (0.5% sulfanilic acid, 0.15% 2-naphthylamine in 30% acetic acid) in a glycine-NaOH buffer. Nitrite ( $\text{NO}_2^-$ ) concentration in the samples was estimated by measuring the absorbance of the red-violet diazo dye at 548 nm.  $\text{NO}_3^-$  concentration in each sample was calculated by subtracting  $\text{NO}_2^-$  concentration value to the total  $\text{NO}_2^-$  concentration, obtained by reducing nitrate ( $\text{NO}_3^-$ ) content in the sample to  $\text{NO}_2^-$  with cadmium. All reactions were conducted in triplicate. Standard  $\text{NO}_2^-$  and  $\text{NO}_3^-$  curves were constructed daily using  $\text{NaNO}_2$  and  $\text{NaNO}_3$  as standard.

To calculate the absolute amount of  $\text{NO}_2^-$  and  $\text{NO}_3^-$ , the concentration in each sample was normalized by the weight of each tissue preparation and collection duration and expressed as  $\mu\text{mol g tissue}^{-1} \text{ min}^{-1}$ . In normal metabolic conditions, the effect of test drugs on  $\text{NO}_2^-$  and  $\text{NO}_3^-$  overflow was expressed as percentage variation with respect to control values. Changes of  $\text{NO}_2^-$  and  $\text{NO}_3^-$  overflow induced by different treatments during *in vitro* I/R, were expressed as percentage variation with respect to values of spontaneous overflow obtained for each treatment in normal metabolic conditions.

## Statistical analysis

For statistical analysis the GraphPad Instat statistical package (version 5.03; GraphPad software, San Diego, CA, USA) was used. Data were analyzed either by one sample *t*-test, Student's *t*-test or analysis of variance (ANOVA) followed by an appropriate *post hoc* comparison tests (Dunnett's or Tukey's Multiple Comparisons tests) or two way ANOVA, as indicated either in the text or in the legends. Differences were considered statistically significant when *P* values  $\leq 0.05$ .

## Drugs and materials

$N^{\omega}$ -[Imino(propylamino)methyl]-L-ornithine hydrochloride ( $N^{\omega}$ -propyl-L-arginine hydrochloride) and 3-aminomethylbenzylacetamide dihydrochloride (1400 W) were purchased from Tocris (Bristol, UK). All other reagents were purchased either from Sigma-Aldrich or from BioRad (Segrate, MI, Italy).

## RESULTS

### Distribution of nNOS-IR and iNOS-IR in LMMP whole-mounts preparations of the guinea pig ileum in normal metabolic conditions and after *in vitro* I/R

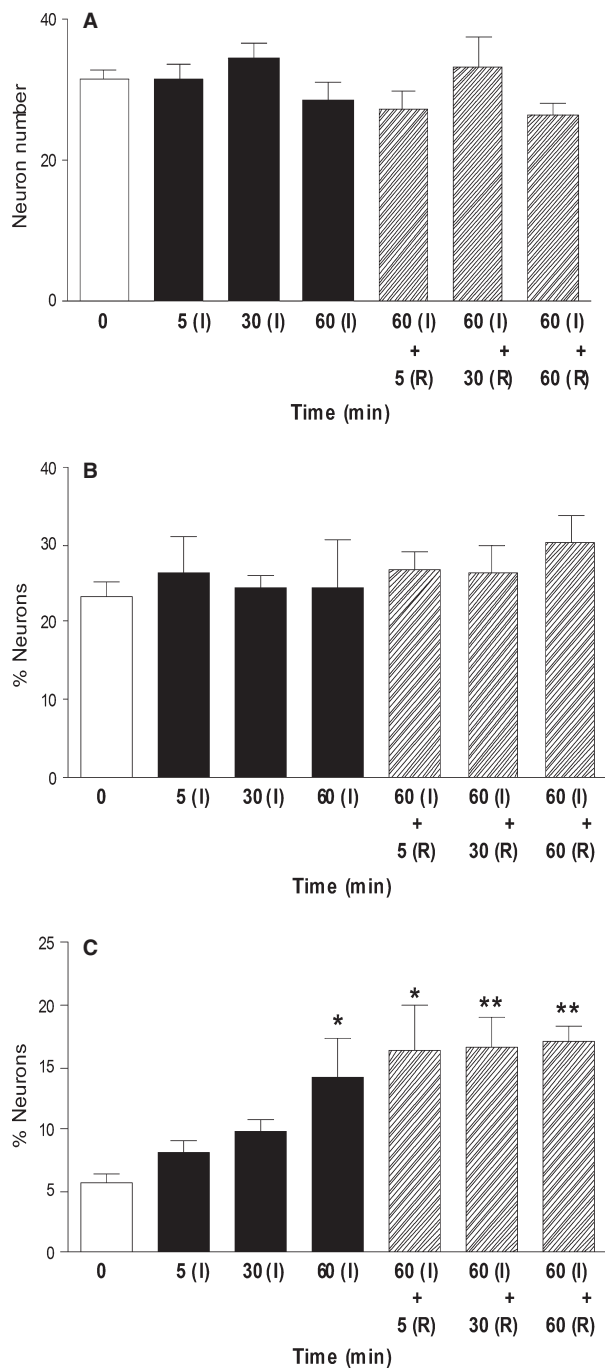
The number of myenteric neurons, as indicated by HuC/D cytoplasmic staining, remained unchanged

during ischemia/reperfusion with respect to control preparations (Fig. 1A). After *in vitro* I/R a proportion of myenteric neurons displayed HuC/D-IR translocation from the cytoplasm to the nucleus (data not shown). In both normal metabolic conditions and after I/R, nNOS-IR was observed in the cytoplasm of myenteric neurons. Some neuronal cells had a large cell body with club-like dendrites, while others were small with an ovoidal shape and with no visible dendrites. nNOS-IR was evidenced also in neuronal fibers within the ganglia and along the interconnecting strands, and in the secondary and tertiary fiber tracts. (Fig. 2A,C). In normal metabolic conditions the percentage of myenteric neurons staining for nNOS was  $23.50 \pm 1.94\%$ ,  $n = 4$ . There was no significant change in the percentage of nNOS immunopositive myenteric neurons at different times after *in vitro* I/R with respect to the value obtained in control preparations (Fig. 1B; Fig. 2D,F, G,I). In both control and I/R preparations, iNOS staining was observed in the soma of myenteric neurons with an ovoidal shape and no visible dendrites (Fig. 3A,C). In normal metabolic conditions the percentage of iNOS immunopositive neurons was  $5.75 \pm 0.75\%$ ,  $n = 4$ . The proportion of iNOS immunopositive neurons significantly increased 60 min after ischemia and during the whole reperfusion period, with respect to the value obtained in controls (Fig. 1C, Fig. 3D,F, G,I). In normal metabolic conditions the majority of iNOS positive neurons ( $97.33 \pm 0.94\%$ ,  $n = 4$ ) expressed nNOS-IR (Fig. 4A–C). All iNOS-IR neurons appearing during I/R conditions were also nNOS immunopositive (Fig. 4D–I). Double labelling with S100 did not reveal the presence of nNOS-IR and iNOS-IR in myenteric glial cells, both in normal metabolic conditions and after I/R (data not shown).

### Levels of expression of nNOS and iNOS in the guinea pig ileum LMMP preparations after *in vitro* I/R

Western blot analysis of nNOS in the guinea pig ileum LMMP preparations (Fig. 5A,B) and in the rat hippocampus (data not shown) revealed one band at 155 kDa. nNOS protein levels in the membrane and soluble fraction of LMMP preparations subjected to I/R conditions remained unchanged with respect to the relative controls (Fig. 5A).

Western blot analysis of iNOS in the guinea pig ileum LMMP preparations (Fig. 5C,D) and in RAW 264.7 macrophage cells (data not shown) revealed one band at 125 kDa. After I/R, iNOS levels significantly increased both in the membrane and soluble fraction of LMMPs, with respect to the relative controls (Fig. 5C).



**Figure 1** (A) Number of myenteric neurons  $\text{mm}^{-2}$  staining for HuC/D in guinea pig ileum whole-mounts preparations in control preparations (empty bar), at different times after inducing *in vitro* ischemia (full bar) and after reperfusion (R, slash). Percentage of myenteric neurons staining for HuC/D and immunopositive for either nNOS (B) or iNOS (C) in guinea pig ileum whole-mounts preparations in control preparations (empty bar), at different times after inducing *in vitro* ischemia (full bar) and after reperfusion (R, slash). Values are expressed as mean  $\pm$  SEM of at least four experiments. Vertical bars indicate SEM \* $P < 0.05$ , \*\* $P < 0.01$  by one way ANOVA followed by Dunnett's test.

nNOS and iNOS protein levels obtained from LMMP preparations superfused with a normal and oxygenated Tyrode's solution for 5, 30, 60, 65, 90 and 120 min after the equilibration period were not significantly different from the amount obtained in control preparations collected at the end of the equilibration period (Fig. 5B,D).

### Levels of nNOS and iNOS mRNA in the guinea pig ileum LMMP preparations after *in vitro* I/R

Quantitative RT-PCR analysis of nNOS and iNOS mRNA in guinea pig ileum LMMP preparations is shown in Fig. 6. Both in control and in I/R preparations analyses of gene expression with respect to  $\beta$ -actin and GAPDH, were not significantly different (data not shown), thus, for the sake of simplification, only  $\beta$ -actin normalized values are reported in Fig. 6.

nNOS mRNA levels significantly increased with respect to the relative controls 60 min after inducing *in vitro* ischemia and after 5 min of reperfusion (Fig. 6A).

iNOS gene expression levels (Fig. 6C) significantly increased 5 min after inducing ischemia and remained elevated throughout the ischemic period. Enhancement of iNOS mRNA levels was also observed during reperfusion (Fig. 6C).

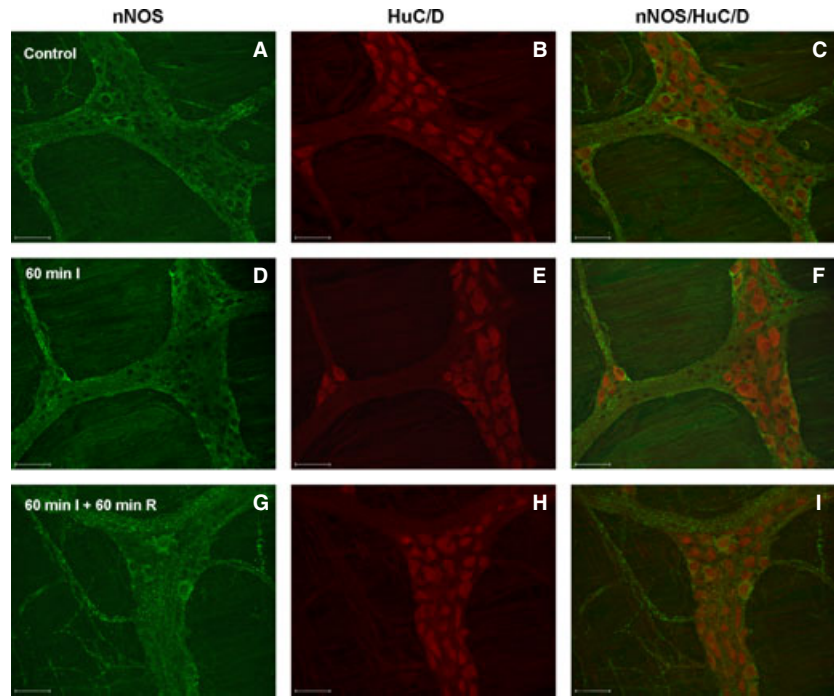
In LMMP preparations superfused with a normal and oxygenated Tyrode's solution for 5, 30, 60, 65, 90 and 120 min after the equilibration period, nNOS and iNOS mRNA levels remained unchanged with respect to control preparations (Fig. 6B,D).

In LMMP control preparations, iNOS levels were about 17-fold lower compared to nNOS levels (data not shown).

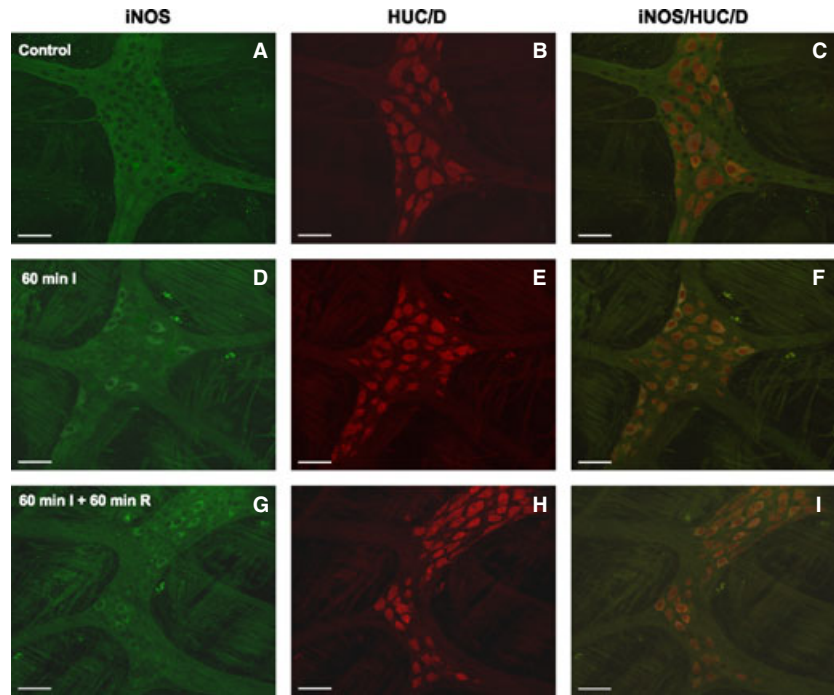
### Effect of *in vitro* I/R on $\text{NO}_2^-$ and $\text{NO}_3^-$ production from the guinea pig ileum

After the 60 min equilibration period,  $\text{NO}_2^-$  and  $\text{NO}_3^-$  concentrations in the superfusate from isolated mucosa-deprived segments of the guinea pig ileum were  $4.02 \pm 0.004 \mu\text{mol g}^{-1} \text{min}^{-1}$  ( $n = 4$ ) and  $11.00 \pm 2.36 \mu\text{mol g}^{-1} \text{min}^{-1}$  ( $n = 4$ ), respectively, and were stable over at least 150 min (Fig. 7A,C).

Both  $\text{NO}_2^-$  and  $\text{NO}_3^-$  concentration significantly increased over control values during the first 5 min after inducing *in vitro* ischemia, and returned to values similar to those obtained in control conditions in the following collection periods under ischemic conditions. A significant enhancement of  $\text{NO}_2^-$  and  $\text{NO}_3^-$  production was observed 5 min after reperfusion.  $\text{NO}_2^-$  and  $\text{NO}_3^-$  overflow tended to reach values not



**Figure 2** Immunohistochemical co-localization of nNOS with the neuronal marker HuC/D in whole-mount preparations of the guinea pig ileum obtained from control preparations (A,B and merged image C), 60 min after *in vitro* ischemia (D,E and merged image F) and at 60 min of reperfusion after 60 min *in vitro* ischemia (G,H and merged image I). The neuronal marker HuC/D stained the somata of all myenteric neurons (B,E,H). Bar 50  $\mu$ m.

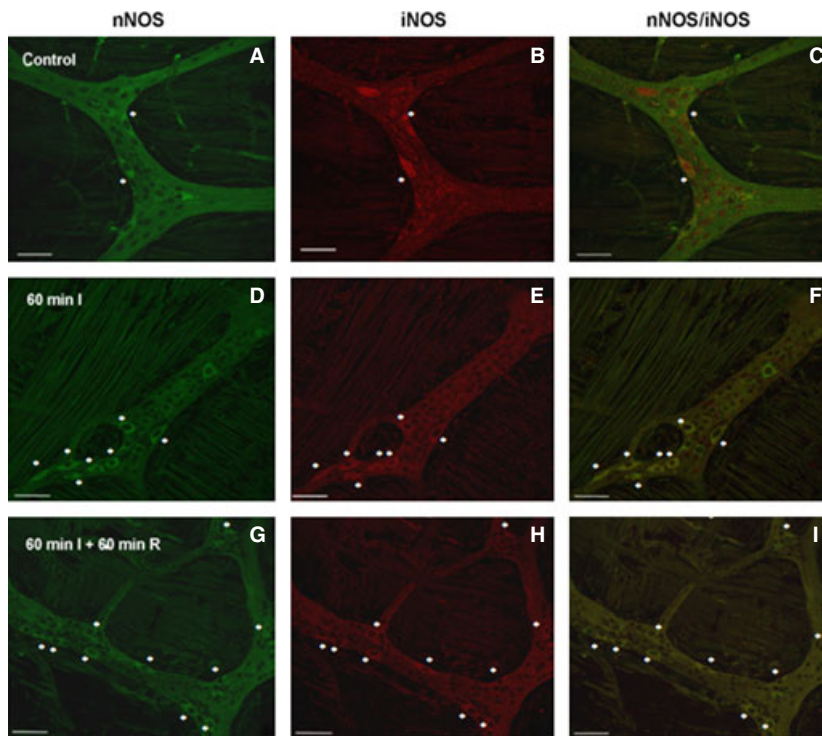


**Figure 3** Immunohistochemical co-localization of iNOS with the neuronal marker HuC/D in whole-mount preparations of the guinea pig ileum obtained from control preparations (A,B and merged image C), 60 min after *in vitro* ischemia (D,E and merged image F) and at 60 min of reperfusion after 60 min *in vitro* ischemia (G,H and merged image I). The neuronal marker HuC/D stained the somata of all myenteric neurons (B,E,H). Bar 50  $\mu$ m.

significantly different from controls 15 min after restoring normal metabolic conditions. The peak of  $\text{NO}_2^-$  and  $\text{NO}_3^-$  concentration observed at 5 min after reperfusion was significantly higher than the values obtained in the last 30 min of *in vitro* ischemia (Fig. 7B,D).

**Effect of nNOS and iNOS blockade on *in vitro* I/R-induced on  $\text{NO}_2^-$  and  $\text{NO}_3^-$  production in the guinea pig ileum**

$N^G$ -propyl-L-arginine (1  $\mu\text{mol L}^{-1}$ ) and 1400 W (10  $\mu\text{mol L}^{-1}$ ) significantly reduced the basal concentrations of



**Figure 4** Confocal images showing co-localization of iNOS and nNOS in myenteric neurons of the guinea pig ileum obtained from control preparations (A,B and merged image C), 60 min after *in vitro* ischemia (D,E and merged image F) and at 60 min of reperfusion after 60 min *in vitro* ischemia (G,H and merged image I). Asterisks indicate myenteric neurons expressing both iNOS and nNOS. Bar 50  $\mu\text{m}$ .

both  $\text{NO}_2^-$  ( $-13.00 \pm 3.91\%$ ,  $n = 7$   $P < 0.05$ ,  $-23.83 \pm 5.21$ ,  $n = 6$   $P < 0.01$ , by one sample *t*-test, respectively) and  $\text{NO}_3^-$  ( $-20.43 \pm 6.73$ ,  $n = 7$   $P < 0.05$ ,  $-30.50 \pm 6.91$ ,  $n = 6$   $P < 0.01$ , by one sample *t*-test, respectively) with respect to control values.

$N^w$ -propyl-L-arginine and 1400 W significantly reduced  $\text{NO}_2^-$  and  $\text{NO}_3^-$  production during I/R. In these conditions, 1400 W reduced  $\text{NO}_3^-$  concentrations to a significantly greater extent than  $N^w$ -propyl-L-arginine (Fig. 8A,B).

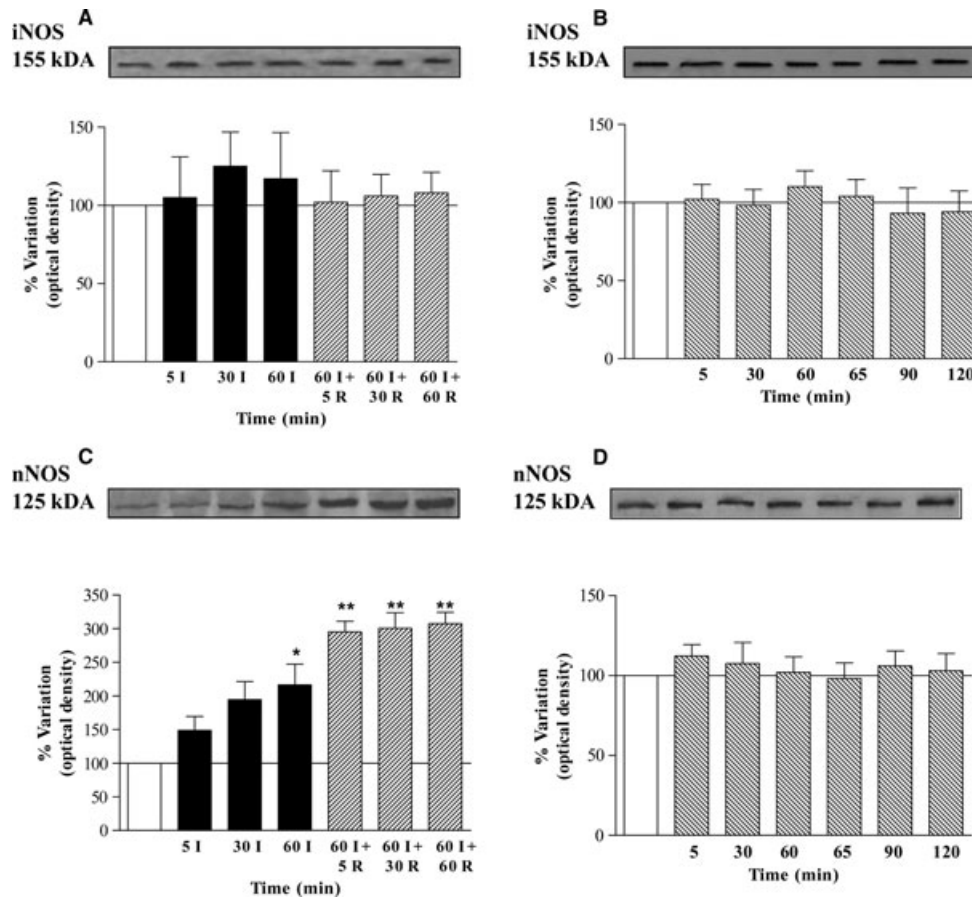
## DISCUSSION

An increasing body of evidence is available to suggest that an I/R injury to the gut may induce severe damage to enteric neurons<sup>6,26</sup> which may involve nitrergic enteric pathways. In the present study we have demonstrated that in the myenteric plexus of the guinea pig ileum, nNOS and iNOS, the predominant sources of NO from enteric neurons in physiological and pathological conditions, respectively,<sup>20,21</sup> may participate to the increased NO production following an acute *in vitro* I/R insult.

Our immunohistochemical findings show that in control preparations approximately 24% of myenteric neurons express nNOS, in agreement with the already reported percentage for nNOS expressing myenteric neurons in the guinea pig, mouse and rat<sup>21,26,32</sup> small

intestine. After *in vitro* I/R in ileal LMMPs, nNOS immunopositive myenteric neurons and protein levels did not change significantly with respect to values obtained under control conditions, while nNOS mRNA levels transiently increased 60 min after ischemia and 5 min after reperfusion. Changes in nNOS mRNA and protein expression have been demonstrated after *in vivo* and *in vitro* I/R injuries both at a central and peripheral level, suggesting that also the constitutive neuronal isoform may play a role during a metabolic insult.<sup>33,34</sup> In human aortic smooth muscle cells, an nNOS promoter has been recently shown to contribute to rapidly up-regulate nNOS mRNA in response to hypoxia with a functionally significant effect on vascular smooth muscle contractility.<sup>35</sup> In our model, nNOS mRNA, but not the protein, is transiently up-regulated. Delay of nNOS mRNA translation into the relevant protein was also demonstrated in the rat nodose ganglion and cerebellum after an hypoxic insult.<sup>33</sup> Interestingly, in the myenteric plexus of the rat ileum, the number of nNOS positive cell bodies significantly increased 24 h after clamping the superior mesenteric artery.<sup>25</sup>

In the present study, immunohistochemical, Western immunoblot and RT-PCR data are provided to suggest that guinea pig ileum myenteric neurons constitutively express iNOS, as already documented by other studies carried out in the mouse, rat and

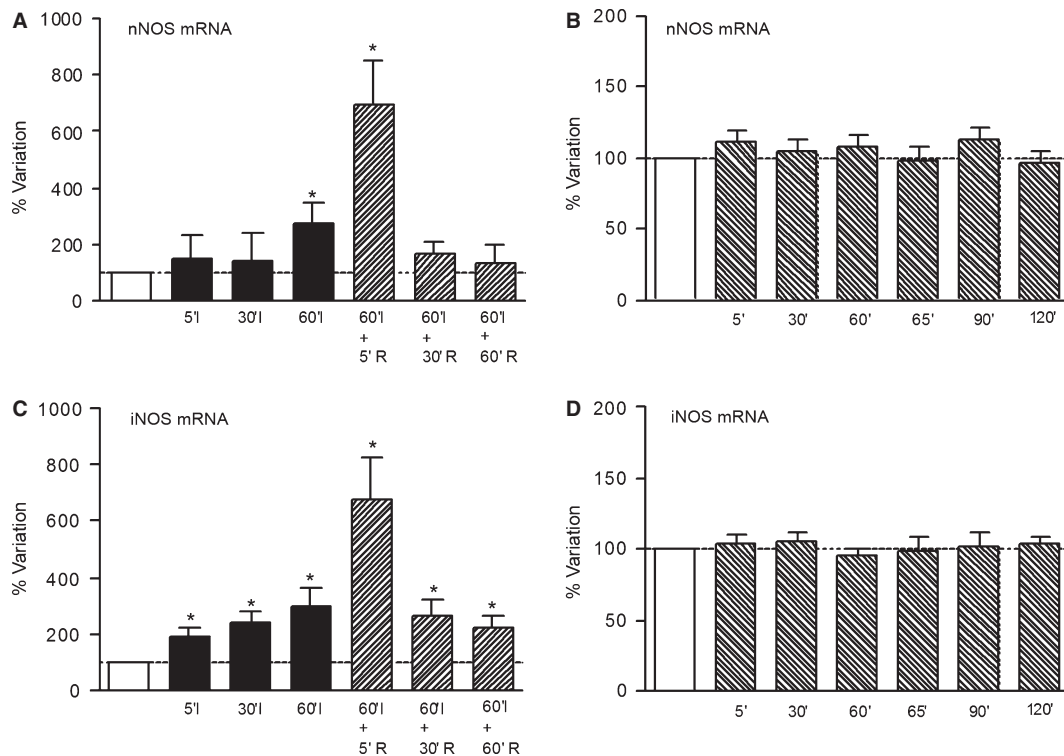


**Figure 5** Levels of expression of nNOS and iNOS in the membrane (P, full bar) and in the supernatant (S, dotted bars) fractions obtained from guinea pig ileum LMMPs preparations at different times after inducing ischemia (I) and reperfusion (R) (A,C) and at different times after the equilibration period in normal metabolic conditions (B,D). Values are expressed as mean  $\pm$  SEM of at least four experiments of the percentage variation of optical density (OD) with respect to relative controls (empty bar). Vertical bars indicate SEM \* $P < 0.05$ , \*\* $P < 0.01$  by one sample *t*-test. Blots representative of immunoreactive bands for nNOS and iNOS in the different experimental conditions are reported on top of each panel. Samples (100  $\mu$ g) were electrophoresed in SDS-10% polyacrylamide gels. Numbers at the margins of the blots indicate relative molecular weights of the respective protein in kDa.

guinea pig gut.<sup>14,17,36</sup> The proportion of myenteric neurons expressing iNOS was smaller (about 6% of the total neuronal population) than the proportion of nNOS expressing neurons. This observation is in agreement with other studies reporting low iNOS immunoreactivity levels in the myenteric plexus of the guinea pig and rat gut.<sup>20,36,37</sup> Accordingly, in control preparations iNOS mRNA levels were significantly lower with respect to those of nNOS. A consistent up-regulation of iNOS has been shown to occur in neuronal cells following I/R.<sup>38,39</sup> Such an adaptive response could be found also in the guinea pig ileum where the percentage of iNOS-expressing myenteric neurons, protein and mRNA levels progressively increased from the first minutes after inducing *in vitro* ischemia and remained elevated during reperfusion. Our data indicate a major involvement of iNOS

rather than nNOS in the changes occurring in myenteric neurons after I/R. In this regard, the observation that, iNOS-IR neurons, appearing after I/R, are nNOS positive may reflect a functional plasticity of myenteric neurons which activate different NOS isoforms depending on either physiological or pathological conditions.<sup>40</sup> In particular, our observations strengthen the concept that in myenteric neurons, nNOS may represent the main source for NO involved in the physiological modulation of non-adrenergic non-cholinergic inhibitory motor responses of the gut. However, under pathological conditions, namely during I/R, within the same cell iNOS may replace nNOS in the synthesis of NO. As regards the molecular mechanisms underpinning upregulation of iNOS expression in our model, one of the most acceptable hypothesis involves hypoxia inducible factor-1 (HIF-1) which is described



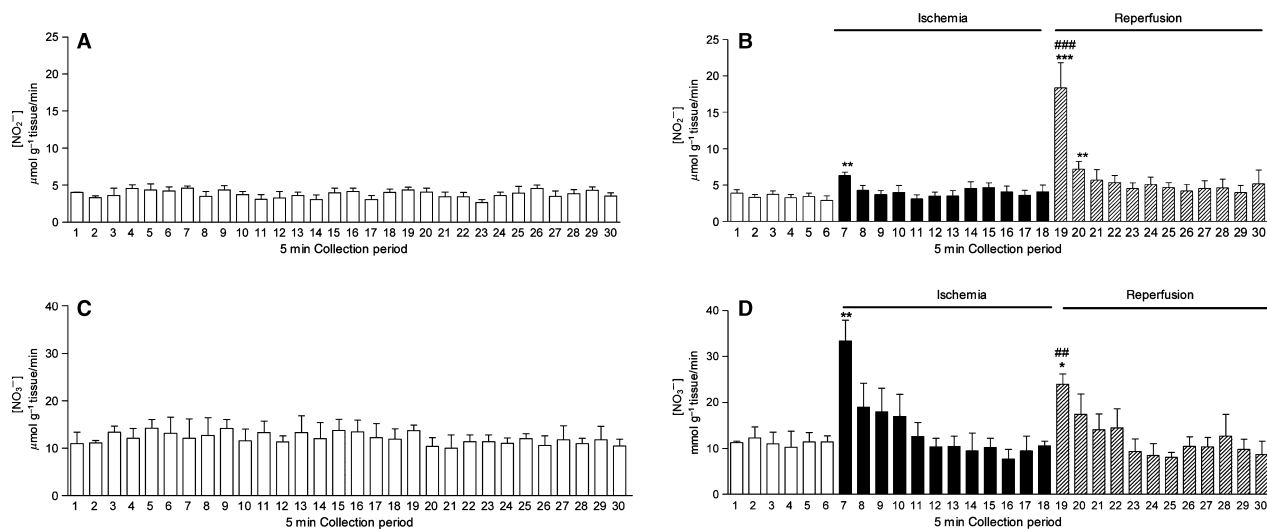


**Figure 6** RT-PCR quantification of nNOS and iNOS transcripts in guinea pig ileum longitudinal muscle myenteric plexus preparations (LMMP) at different times after inducing *in vitro* ischemia (I, full bar), reperfusion (R, slash) (A,C) and at different times after the equilibration period in normal metabolic conditions (backslash) (B,D). Values are mean  $\pm$  SEM of at least four experiments of the percentage variation of relative gene expression with respect to relative controls (empty bar). Vertical bars indicate SEM. \*  $P < 0.05$  by one sample *t*-test. The relative gene expression was determined by comparing  $\Delta\Delta CT$  values normalized to  $\beta$ -actin.

to mediate iNOS mRNA transcription under ischemic/reperfusion conditions.<sup>41</sup>

In the guinea pig ileum, the concentrations of NO oxidation products, namely  $\text{NO}_2^-$  and  $\text{NO}_3^-$ , transiently increased at the onset of hypoxia/hypoglycaemia and after returning to an oxygenated glucose-containing medium. This finding suggests that the first minutes after inducing *in vitro* ischemia and reperfusion are crucial moments for the modulation of NO synthesis in our model. NO is a reactive and unstable molecule and its half-life is estimated as a few seconds in biological specimens.<sup>42</sup> Thus, to assess its synthesis in our preparations we have quantitatively determined the concentrations of its stable oxidation products,  $\text{NO}_2^-$  and  $\text{NO}_3^-$  resorting to a reproducible and sensitive spectrophotometric method assay based on the Griess reaction.<sup>43</sup> In biological specimens,  $\text{NO}_3^-$  concentrations are in the micromolar range and are 10–100-fold higher than  $\text{NO}_2^-$ , since this latter NO oxidation product is almost completely oxidized to  $\text{NO}_3^-$  by oxyhaemoglobin or oxymyoglobin.<sup>44</sup> In agreement with the literature,  $\text{NO}_3^-$  concentrations from ileal superfusate resulted in the micromolar range, although this value was just three fold higher than

$\text{NO}_2^-$  concentrations. Higher  $\text{NO}_2^-$  levels may be explained considering that in the superfusing medium, consisting of an aqueous solution lacking of haemoproteins,  $\text{NO}_2^-$  are more stable and represent a conspicuous fraction of spontaneous NO oxidation products, as already suggested for the rabbit stomach and colon.<sup>45</sup> Under normal metabolic conditions, the ability of  $N^{\omega}$ -propyl-L-arginine and 1400 W, to decrease  $\text{NO}_2^-$  and  $\text{NO}_3^-$  levels is suggestive of a spontaneous NO release from the guinea pig ileum, which involves both nNOS and iNOS activation. Spontaneous NO synthesis has been demonstrated from the canine proximal colon,<sup>46</sup> and may have a role in the inhibition of intestinal smooth muscle spontaneous excitability.<sup>47</sup> Sensitivity to 1400 W strengthens the hypothesis of the presence of a constitutively active functional iNOS in the guinea pig ileum, as already suggested for the human colon and mouse ileum.<sup>21,48</sup> Our immunohistochemical data suggest that, in our preparations, myenteric neurons may represent the major cellular population releasing NO, via iNOS activation. Although we could not find evidence of either nNOS-IR or iNOS-IR in enteric glial cells by immunohistochemistry, we cannot entirely exclude the



**Figure 7**  $\text{NO}_2^-$  and  $\text{NO}_3^-$  concentrations in the superfusate from isolated segments of the guinea pig ileum in normal (A,C), in *in vitro* ischemic conditions and during reperfusion after *in vitro* ischemia (B,D). Horizontal bars indicate the period of treatment. Each column is the mean of at least 4–7 experiments. Vertical bars indicate SEM, \* $P < 0.05$ , \*\* $P < 0.01$ , \*\*\* $P < 0.001$  (vs controls) and ### $P < 0.01$ , #### $P < 0.01$  vs last six collections in ischemic conditions by one way ANOVA followed by Tuckey's *post hoc* test.

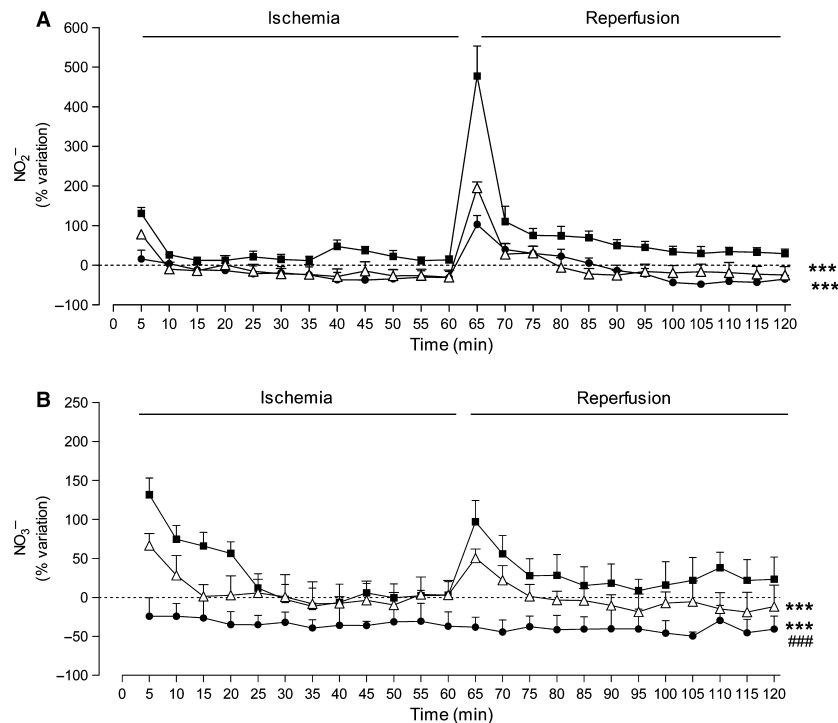
participation of this cell population. Indeed, in astrocytes, which share many common morphological and functional features with enteric glial cells, iNOS represents a major source for NO under physiological conditions.<sup>40,49</sup>

After inducing both *in vitro* ischemia and reperfusion NO ileal levels rapidly and transiently increased via a NOS dependent pathway, involving both nNOS and iNOS, as suggested by the ability of *N*<sup>o</sup>-propyl-L-arginine and 1400 W to reduce the peak concentrations of both  $\text{NO}_2^-$  and  $\text{NO}_3^-$ . The higher sensitivity of  $\text{NO}_3^-$  levels to 1400 W, with respect to *N*<sup>o</sup>-propyl-L-arginine, may be indicative of a major role played by iNOS in myenteric neurons during pathological conditions, as expected. However, during the first min after reperfusion, NOS may not represent the unique source for NO synthesis, as indicated by the partial ability of NOS inhibitors to decrease  $\text{NO}_2^-$  levels, in these conditions. In this regard, we may hypothesize that a fraction of  $\text{NO}_2^-$  may originate from NO bound to scavengers, such as deoxygenated haemoglobin and myoglobin.<sup>50,51</sup> Under hypoxic conditions, deoxyhaemoglobin and deoxymyoglobin may mediate the non-enzymatic conversion of  $\text{NO}_2^-$  to NO, representing sources of NO.<sup>52</sup> Additionally, we cannot exclude that, in the guinea pig ileum, NO may be stored in the form of molecular complexes, such as S-nitrosothiols and dinitrosyl iron complexes, from which NO can be subsequently released.<sup>53</sup>

In our model the relationship between NO production and changes of NOS expression is not consistent throughout the experiment, as already observed in the

human colon from IBD patients.<sup>21</sup> Indeed, the question whether an increase of NOS expression may lead to higher NO production is controversial. Recently in the rat myocardium elevation of NO levels during re-oxygenation after *in vitro* hypoxia was transitory and did not reflect the long lasting changes of both iNOS and eNOS expression and activity.<sup>54</sup> In the guinea pig ileum, enhancement of NO synthesis during the first min after ischemia and reperfusion was paralleled by an increase of iNOS expression. However, nNOS protein levels remained unchanged suggesting that the possible enhancement of nNOS activity may depend upon post-translational events modulating the enzyme function.<sup>34</sup> One of the possible mechanisms may be represented by a  $\text{Ca}^{++}$ -dependent nNOS activation consequent to glutamate mediated overstimulation of myenteric NMDA receptors during the I/R insult.<sup>27</sup> Reduction of NO levels during maintained *in vitro* ischemic conditions, in spite of the increased iNOS expression and the unchanged nNOS expression, may depend upon hypoxia-mediated inhibition of NOS activity.<sup>41,55</sup> Furthermore, NO binding to scavenger molecules in the ileal tissue during both *in vitro* ischemia and reperfusion, may reduce the concentration of NO oxidized species in the superfusate.<sup>52</sup> During reoxygenation of the ischemic tissue, NO may be captured by molecular components of the tissue and converted into reactive nitrogen species which contribute to protein nitrosative modifications, cellular apoptosis and necrosis.<sup>22,56</sup>

From a functional viewpoint, the boost of NO synthesis after an acute I/R damage may be the basis



**Figure 8** Effect of *N*<sup>o</sup>-propyl-L-arginine (1  $\mu\text{mol L}^{-1}$ ) and 1400 W (10  $\mu\text{mol L}^{-1}$ ) on  $\text{NO}_2^-$  (A) and  $\text{NO}_3^-$  (B) production from the guinea pig ileum evoked by *in vitro* ischemia and reperfusion. Horizontal bars indicate the period of treatment. *In vitro* ischemia/reperfusion only (■) *N*<sup>o</sup>-propyl-L-arginine ( $\Delta$ ) and 1400 W (●). Data are expressed as percentage variation with respect to values of spontaneous overflow obtained for each treatment in normal metabolic conditions. Each point represents the mean of 4–7 experiments. Vertical bars indicate SEM; \*\*\**P* < 0.001, by two way ANOVA vs *in vitro* ischemia/reperfusion only. ###*P* < 0.001 vs *N*<sup>o</sup>-propyl-L-arginine-treated preparations.

of the reduced muscular contractility described soon after an hypoxic injury to the intestine.<sup>10</sup> Enhancement of NO synthesis may also retain an emergency function, exerting a vasodilator effect in order to regulate the blood flow through the hypoxic tissues.<sup>55</sup> In addition, immediate iNOS expression changes, which are not reversed by reperfusion, and potential delayed alterations of nNOS protein may contribute to NO-mediated myenteric neuron damage underlying post-ischemic impairment of intestinal transit.<sup>6,23,25</sup>

In conclusion the present study has provided evidence that in the guinea pig ileum during *in vitro* I/R, both iNOS and nNOS are involved in acute overproduction of NO. Changes of iNOS protein, and possibly of nNOS protein, may participate to the delay of the intestinal transit and foster long lasting effects on enteric neuronal circuitries in these conditions. A more clear-cut differentiation of the function of different NOS isoforms in clinical conditions, such as during an ischemic gastrointestinal injury, may provide useful information about the pathophysiological basis of the disease and eventually lead to the devel-

opment of pharmacological strategies targeting specific NOS.

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## DISCLOSURE

The authors state no potential conflicts of interest.

## AUTHOR CONTRIBUTION

GC analysed data, wrote the paper; GC EC, SM, RO EM, BB and DV designed the research study and performed the research; GF, SL and FC contributed to design the research and to discuss data.

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