Urinary N-nitrosoproline excretion: a further evaluation of the nitrosamine hypothesis of gastric carcinogenesis in precancerous conditions

C N HALL, J S KIRKHAM, AND T C NORTHFIELD

From the Norman Tanner Gastroenterology Unit, St James' Hospital, Balham, London

SUMMARY Measurement of N-nitroso compounds in gastric juice by different methods has given conflicting results. In order to resolve this controversy, we have assessed endogenous nitrosation by the independent N-nitrosoproline excretion test in subjects who had previously undergone gastric juice analysis by one of these methods. Ten Polya gastrectomy, 10 pernicious anaemia and nine matched control subjects were fed 380 mg of nitrate in beetroot juice and 500 mg proline. N-nitrosoproline (N-Pro) synthesised intragastrically from these precursors, and quantitatively excreted by the kidneys, was measured in 24 hour urine samples (collection checked by creatinine clearance). N-Pro excretion (mean±SEM) was reduced (p<0.01) in pernicious anaemia (1.1±0.8 ng/day) compared with matched control (18.0±7.2 ng/day), and also tended to be lower (NS) in polya gastrectomy (3.2±2.3 ng/day). Twenty four hour intragastric pH was monitored on a separate occasion in 23 of the 29 subjects; 13 were hypoacidaic (pH>4.50% of 24 hours) and 10 were acidic. N-Pro yields were reduced (p<0.01) in the hypoacidaic group (0.9±0.6 ng/day) compared with the acidic group (17.9±6.6 ng/day), and N-Pro was negatively associated with mean intragastric pH (\(\tau = -0.53, p=0.001\)). We conclude that endogenous synthesis of this specific N-nitroso compound is favoured by low rather than high pH. These results are concordant with those previously reported in gastric juice from the same subjects and suggest that nitrosation is chemically rather than bacterially mediated, contrary to the nitrosamine hypothesis of gastric carcinogenesis.

The nitrosamine hypothesis of gastric carcinogenesis postulates that high intragastric pH in hypochlorhydria promotes the growth of bacteria which reduce dietary nitrate to nitrite and then convert dietary amines, in the presence of this nitrite, into carcinogenic N-nitroso compounds.\(^1\) This hypothesis has been tested by the measurement of N-nitroso compound concentrations in gastric aspirate from subjects with medically and surgically induced hypochlorhydria and in controls, but the results are conflicting. Increased concentrations have been reported as being associated with high pH\(^1\) or as being unrelated.\(^5\) These discrepancies may be accounted for by differences in methodology, for a wholly satisfactory method for the determination of N-nitroso compounds in complex biological fluids is as yet unavailable. An indirect method for the assessment of endogenous nitrosation has recently been described,\(^10\) and this obviates the need for gastric juice analysis. Instead, N-nitrosoproline excreted in the urine after the ingestion of precursors provides a quantitative measurement of nitrosation of ingested proline in gastric juice. We therefore carried out this test on Polya gastrectomy, pernicious anaemia and matched control subjects and compared the results with data on total N-nitroso compounds in gastric juice obtained in a previous study of these same individuals.

Address for correspondence: Dr T C Northfield, St Georges Hospital Medical School, Dept of Medicine II, Blackshaw Road, London SW17 0RE.

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Methods

Subjects
Ten Polya gastrectomy subjects (Visick I and II) operated on at least 15 years earlier for duodenal ulcer were compared with nine matched controls comprising healthy volunteers free of gastrointestinal disease, and with 10 pernicious anaemia subjects characterised by a previous diagnosis of macrocytic anaemia, vitamin B12 deficiency and vitamin B12 malabsorption correctable by intrinsic factor. Subjects were matched in nine triplets (plus an additional Polya gastrectomy-pernicious anaemia pairing) for race, sex, and age to the nearest decade (Table). Matching of groups was also good for height, but pernicious anaemia tended to be heavier than the other two groups (NS).

On the day of study, fasting subjects voided urine and provided a blood sample before being fed 380 mg nitrate in beetroot juice (200 ml) followed 30 minutes later by 500 mg of proline in water (100 ml). After another four hour fast (during which time smoking was not permitted), a standardised solid meal was given and then a normal daily routine was adopted. During this latter phase, subjects were merely advised to avoid potential sources of preformed N-Pro such as cured meat products and beer. Urine was collected for 24 hours in receptacles containing 10 mg sulphamic acid (to prevent artefactual formation of N-Pro). Creatinine was measured in serum and fresh urine and creatinine clearance used as an approximate indicator of completeness of urine collection. Thirty millilitre aliquots of urine were stored at −20°C before analysis.

N-nitrosoproline was extracted from urine according to the method of Sen and Seaman. Twenty micrograms of N-pipercolic acid (internal standard) was added to 15 ml urine followed by 3 ml of 3N sulphuric acid containing 1% sulphamic acid and then the mixture placed on a Clin Elut Extube (Analyticchem International, Harbour City, CA, USA) and allowed to equilibrate for five minutes. N-nitrosamino acids were eluted with 4×20 ml of ethyl acetate, waiting two to three minutes between each elution. The eluate was dried for 30 minutes over anhydrous sodium sulphate, filtered and reduced to 2 ml using a rotary evaporator. The resulting solution was transferred (with adequate rinsing with ethyl acetate) to a 15 ml graduated test tube and concentrated to 0-1 ml (avoiding evaporation to dryness) in a stream of nitrogen. Finally 1 ml BF3-methanol reagent was added, the test tube stoppered and this mixture heated in a sand bath at 65°C for 30 minutes. Upon cooling to room temperature, 4 ml water and exactly 1 ml dichloromethane were added before agitating in a vortex mixer for two minutes. N-nitrosamino acids were then determined by placing 6 μl of the dichloromethane layer into a gas-liquid chromatograph fitted with a thermal energy analyser. Mean recovery (±SEM) of internal standard was 65±2%.

Statistical Analysis
Matched data were evaluated by paired Student's t test and unmatched data by Wilcoxon's rank sum test. The correlation between urinary N-Pro and mean intragastric pH was analysed by Kendall's rank correlation coefficient because of the many negative yields of N-Pro and the non-normal distribution of these data.

Table

<table>
<thead>
<tr>
<th></th>
<th>Polyga gastrectomy</th>
<th>Control</th>
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</tr>
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<tbody>
<tr>
<td>Age (yr)</td>
<td>65±3</td>
<td>64±3</td>
<td>67±3</td>
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<td>Sex (M:F)</td>
<td>7:3</td>
<td>6:3</td>
<td>7:3</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>171±4</td>
<td>170±3</td>
<td>172±4</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>66±4</td>
<td>65±4</td>
<td>72±5</td>
</tr>
<tr>
<td>Race</td>
<td>All caucasian</td>
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</tr>
</tbody>
</table>

Fig. 1 Assessment of completeness of 24 hour urine collection. (PG = Polya gastrectomy, PA = pernicious anaemia)
Results

Creatinine clearance was similar in all subjects, and there were no significant differences in clearance between groups, indicating that urine collection was very likely to have been complete (Fig. 1). N-nitrosoproline was detected in the urine of all but one matched control but in only three Polya gastrectomy and two pernicious anaemia subjects; consequently excretion (mean±SEM) was reduced (p<0.01) in pernicious anaemia (1.1±0.8 ng/day) compared with matched controls (18.0±7.2 ng/day) and also tended to be lower (NS) in Polya gastrectomy (3.2±2.3 ng/day) (Fig. 2).

Twenty four hour intragastric pH profiles were available for 23 of the 29 subjects because of their recent participation in a study of N-nitroso compounds in gastric juice. Thirteen of these individuals were defined as hypoacidic (pH>4>50% of 24 hours) and 10 were acidic. Only two hypoacidic and all but one acidic individuals produced N-Pro and excretion (mean±SEM) was reduced (p<0.01) in hypoacidic (0.9±0.6 ng/day) compared with acidic subjects (17.9±6.6 ng/day) (Fig. 3). N-nitrosoproline was invariably present in the urine of subjects with a mean intragastric pH<4 and only rarely detected above this level. In addition, N-Pro yields were negatively associated with mean intragastric pH (tau=−0.53; p=0.001) (Fig. 4).

Discussion

In order to evaluate the N-nitroso compound hypothesis of gastric carcinogenesis, direct measurement of total N-nitroso compound concentrations in aspirated gastric juice has been carried out, but this work is hampered by the unavailability of a wholly satisfactory method for the determination of these agents in complex biological fluids. Estimations have been done either on single samples of fasting gastric juice taken at endoscopy and stored prior to analysis or on multiple samples taken over 24 hours and analysed immediately upon retrieval. While single sample techniques may not fully represent intragastric events and analysis of stored material is liable to underestimate total compounds because of loss of volatile and unstable species, the sole method of immediate estimation may not be entirely selective and may therefore be prone to overestimation. In general, total N-nitroso compound concentrations measured in fresh samples are approximately five to 10-fold greater than in stored material. Furthermore, the reported relationship of total N-nitroso compounds to intragastric pH is

Fig. 2  Urinary yields of N-nitrosoproline in Polya gastrectomy (PG), pernicious anaemia (PA) and matched controls (MC).

Fig. 3  Urinary yields of N-nitrosoproline in acidic and hypoacidic groups.
bacterial mechanisms enhance nitrosation at high intragastric pH. Our results are consistent in showing that both total N-nitroso compounds in gastric juice and N-nitrosoproline in urine are reduced in hypoxic subjects, and this suggests that chemical mechanisms of nitrosation predominate in vivo in man contrary to the N-nitroso compound hypothesis.

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