Phase I study of intra-arterial interleukin-2 in squamous cell carcinoma of the head and neck

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Interleukin-2 (IL-2) administered intravenously as a single agent or together with lymphokine activated killer cells has been shown to have activity against a variety of tumours with the highest response rates recorded in patients with renal cell carcinoma and melanoma (21–35%, Rosenberg et al., 1989). The precise mechanism by which IL-2 causes tumour regression is uncertain but it may involve the activation of cellular immune mechanisms. In vivo IL-2 stimulates the proliferation of a variety of lymphoid cells including activated T cells, natural killer cells, lymphokine activated killer cells, B cells and macrophages (Gillis & Smith, 1977; Henny et al., 1981; Grimm et al., 1982; Waldmann et al., 1984; Malkovsky et al., 1987). IL-2 is a powerful regulator of the immune system and although it is detectable in the serum of healthy controls and cancer patients (Lissoni et al., 1990) its main site of action is probably at a local level. Therefore, a more physiological approach to the administration of IL-2 might be to deliver it locally to sites where potentially tumoricidal lymphocytes may be concentrated. Such loco-regional therapy should also be associated with fewer systemic side effects. Continuous intralymphatic infusions over a period of days are not technically possible but a similar result could be achieved by the intra-arterial route. Patients with tumours of the head and neck were selected for this treatment approach as the arterial supply to these tumours is frequently accessible to cannulation. In addition these lesions are often easily biopsied and provide a unique opportunity to study the histopathological and immunohistochemical changes associated with the local delivery of varying concentrations of IL-2.

We present the first stage of this programme which is a phase I dose escalation study of intra-arterial IL-2 administered by continuous infusion over a maximum of 10 days to patients with incurable squamous cell carcinoma of the head and neck.

Materials and methods

Patients

Patients with recurrent or untreated, but incurable squamous cell carcinoma of the head and neck were eligible for the study. Patients had to satisfy the following criteria before entry: performance status 0–1 (ECOG), WBC > 4 x 10⁹ per litre, platelets > 100,000 x 10⁹ per litre, HCT > 30%, normal serum bilirubin and creatinine and the principal blood supply to the tumour had to be from a branch of the external carotid artery. Patients were excluded if they had a significant history of cardiovascular disease, a contra-indication to the use of pressor agents, a previous organ allograft, a serious active infection, a requirement for corticosteroids or had a concurrent second primary malignancy. Patients gave fully informed, witnessed, written consent as laid down by the Ethics Committee of the Royal Marsden Hospital, London.

A total of 15 patients fulfilled the entry criteria but two refused treatment and one was unable to give fully informed consent for psychological reasons. Thus, 12 patients (nine male, three female) satisfied the final entry criteria. The median age of patients was 55 years (range 31–72) and the majority (nine) had head and neck cancer (carcinoma of the head and neck). Two patients had recurrent disease after radiotherapy and/or surgery and two had previously untreated but incurable disease.

Administration

High-flow nylon (4.0 G) or vertebral (3.7 G; 4.7 G) catheters (William Cook Europe Ltd) were inserted retrogradely under general anaesthetic via an arteriotomy in the superficial temporal artery in all but two cases. In these patients the catheters were inserted directly into the maxillary and superior thyroid arteries. The position of the catheter was checked by angiography or by the instillation of fluorescein (10% w/v) into the line. Heparin (50 iu in 50 ml normal saline) was infused at 2.5 ml h⁻¹ through the catheter for 24 h while patients recovered from the general anaesthetic. Two out of the first three patients entered developed line blockages/arterial thrombosis during the first 48 h of treatment and therefore all subsequent patients were systemically anti-coagulated with intravenous heparin. The dose was altered daily according to the partial thromboplastin time. IL-2 in 60 ml of 5% dextrose with 2.5% albumin (final concentration) was administered daily over 24 h by continuous intra-arterial infusion via a syringe pump.

Study design

We intended to enter three patients at each dose level which were as follows: level 1, 3 x 10⁴ iu/day; level 2, 3 x 10⁵ iu/day; level 3, 3 x 10⁶ iu/day; level 4, 3 x 10⁷ iu/day (Table I). The first patient was treated for 5 days, patients 2–4 for 10 days with a 2-day break after 5 days and subsequently it was planned that all patients should receive continuous treatment for 10 days. However, the two patients treated at the highest dose level (3 x 10⁷ iu/day) required a 2-day break after 5 days of treatment because of systemic toxicity.

The end point of the study was the development of systemic toxicity typical of intravenous interleukin-2 at standard dose which for most continuous infusion regimens is 18 x 10⁶ iu m² day.

Table I Dose schedule

| Level | Dose iu/(24 h) | Schedule days | n |
|-------|---------------|---------------|---|
| 1     | 3 x 10⁴       | 5             | 1 |
| 2     | 3 x 10⁵       | 5-5-5         | 2 |
| 3     | 3 x 10⁶       | 10            | 3 |
| 4     | 3 x 10⁷       | 5-5-5         | 2 |

*Equivalent iv treatment dose.

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Assessment of toxicity and response

Patients were assessed daily for subjective toxicity (WHO criteria) and for local complications. For those toxicities where no WHO grade exists an arbitrary scale was used, side effects were recorded as mild (grade 1), moderate (grade 2), severe (grade 3) or life threatening (grade 4). Venepuncture was performed daily to measure FBC, serum creatinine, urea and electrolytes, clotting studies, liver function tests, gamma GT, albumin and calcium. Chest X-ray, blood cultures and MSUs were performed weekly. Pre- and post-treatment biopsies were obtained in 11 patients; one patient had a pre-treatment biopsy only as she refused biopsy post-treatment.

Tumour response was defined according to standard criteria: Complete response (CR) was defined as the disappearance of all clinical, radiological and biochemical evidence of disease for at least 1 month; partial response (PR) was defined as a reduction in the product of two diameters of measurable disease by at least 50% for at least 1 month (Miller et al., 1981).

Results

Patients treated at dose levels 1–3 did not experience any significant systemic side effects (Tables II and III) and remained fully mobile and capable of self-care. However systemic toxicity typical of iv IL-2 at standard dose was seen at the highest dose level (level 4; 3 × 10^5 IU day). The two patients treated at this dose experienced anorexia, nausea, fever, fatigue, weight gain and transient abnormalities of liver function (grade 1). In addition, one of these patients became hypocalbuminaemic and experienced episodes of hypotension (grade 3) while the other complained of shortness of breath (grade 2) and dry desquamation of his skin (grade 2).

The commonest toxicities encountered were local: tumour pain, eight patients; facial oedema, eight patients; infection/facial cellulitis, four patients. These side effects were not dose-related and resolved during the week following cessation of treatment. The four patients with signs of local cellulitis were treated with antibiotics although none had a positive blood culture. Two patients had haemorrhages from their tumours that may have been related to their anticoagulation. Initially there were considerable problems maintaining the patency of the arterial lines. The first three patients were not systemically anticoagulated and two of these patients had line failures within 48 h of catheter insertion due to arterial thrombosis and the lines were consequently removed. One of these patients was able to complete treatment after the catheter was resited. There were no obvious local factors to account for these failures and once subsequent patients were systemically anticoagulated this problem did not recur.

Seven patients developed eight positive bacterial cultures at the following sites: sputum, three cultures (Candida albicans × two patients, B. haemolytic streptococcus); arterial catheter tip, two cultures (Klebsiella pneumoniae, Staphylococcus epidermidis); wound, two cultures (Haemolytic streptococcus, Staphylococcus epidermidis); blood, one culture (Staphylococcus epidermidis).

Two patients who received the lowest dose of IL-2, 3 × 10^4 iv day had partial responses and no response were seen at doses above this level.

| Table II | Subjective toxicity. Worse WHO grade recorded at all dose levels |
|----------|---------------------------------------------------------------|
| WHO Grade | 0 | 1 | 2 | 3 | 4 |
| Fever    | 2 | 2 | 8 | 0 | 0 |
| Wt loss  | 8 | 0 | 4 | 0 | 0 |
| Oedema   | 9 | 1 | 2 | 0 | 0 |
| CNS      | 6 | 5 | 1 | 0 | 0 |
| Sore mouth | 6 | 5 | 1 | 0 | 0 |
| Nausea/Vomiting | 6 | 5 | 1 | 0 | 0 |
| Anorexia | 7 | 3 | 1 | 0 | 0 |
| Weight loss | 9 | 0 | 3 | 0 | 0 |
| SOB      | 8 | 3 | 1 | 0 | 0 |
| Pruritus | 9 | 2 | 1 | 0 | 0 |
| Diarrhoea | 8 | 4 | 0 | 0 | 0 |
| Taste    | 10 | 2 | 0 | 0 | 0 |
| Rigor    | 11 | 1 | 0 | 0 | 0 |

CNS = central neurological symptoms. SOB = shortness of breath.

| Table III | Objective toxicity. Worse WHO grade recorded at all dose levels |
|-----------|---------------------------------------------------------------|
| WHO Grade | 0 | 1 | 2 | 3 | 4 |
| Anaemia   | 6 | 2 | 3 | 1 | 0 |
| Hypotension | 8 | 0 | 3 | 1 | 0 |
| Infection | 8 | 0 | 3 | 1 | 0 |
| Liver     | 8 | 3 | 1 | 0 | 0 |
| Proteinuria | 8 | 3 | 1 | 0 | 0 |
| Platelets | 11 | 1 | 0 | 0 | 0 |

Discussion

We have found that significant systemic side effects do not occur with intra-arterial IL-2 at doses of 3 × 10^5 IU day and below. At a dose of 3 × 10^5 IU day systemic toxicity is similar to that seen with standard intravenous regimen. Local complications were greater than expected particularly those of cellulitis, facial oedema and arterial thrombosis. The cellulitis and local oedema were not due to infection although we and others have reported an increased risk of infection associated with IL-2 therapy (Hartmann et al., 1989; Bock et al., 1990; Hardy et al., 1990). Histological examination of post-treatment biopsies did not reveal any specific explanation for this complication (data not shown). Arterial thrombosis was a major problem during the early part of the study until patients were systemically anti-coagulated. There have been previous reports of local thrombus associated with intra-arterial IL-2 (Klasa & Silver, 1989; Eggermont et al., 1990; Mavligit et al., 1990) and it has been suggested that this is due to direct damage to the vascular endothelium by IL-2 activated lymphocytes (Dame et al., 1987), activation of the intrinsic system of coagulation (Fleischmann et al., 1991) or an increase in the coagulant properties of endothelium by IL-2 induced cytokines such as IL-1 and tumour necrosis factor (Cotran & Pober, 1989).

Phase II trials in patients with squamous cell carcinoma of the head and neck utilising perilymphatic injections have been performed but reported response rates vary widely, from 0–8% (Selvaghi et al., 1990; de Mulder et al., 1989) to 65% (Cortesina et al., 1991). All these studies suggested that systemic toxicity is absent when this approach is employed, but at doses of 10^6 IU local swelling and pain occurred similar to that observed in our patients (Cortesina et al., 1991). In these studies bolus doses were given and thus the tumour infiltrating lymphocytes were only intermittently exposed to IL-2. It is possible that continuous exposure to IL-2 might result in a greater anti-tumour effect.

Head and neck cancers are often easily accessible to biopsy and therefore a programme of locally infused IL-2 presents a unique opportunity to study the precise changes that take place within tumours as a result of prolonged exposure to both high and low local concentrations of IL-2. In addition, intra-arterial IL-2 studies have important implications for those groups who are studying targeted gene therapy. Unless continuous intra-tumoral infusions of IL-2 cause tumour regressions then it is unlikely that a strategy utilising IL-2 secreting tumour infiltrating lymphocytes will be of benefit in this context. We suggest that groups currently working on targeted gene therapy study the effects of locally infused IL-2 before embarking on any clinical trials.
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