Multivariate regression analyses of data from a randomised, double-blind, placebo-controlled study confirm quality of life benefit of epoetin alfa in patients receiving non-platinum chemotherapy

L Fallowfield*,1, D Gagnon2, M Zagari3, D Cell4, B Bresnahan5, TJ Littlewood6, P McNulty7, G Gorzegno6 and M Freund7 for the Epoetin Alfa Study Group8

1Cancer Research UK Psychosocial Oncology Group, Brighton and Sussex Medical School, University of Sussex, Brighton, BN1 9QG, UK; 2Johnson & Johnson Pharmaceutical Research & Development, L.L.C., 700 Route 202, Raritan, New Jersey, NJ 08869, USA; 3Ortho Biotech UK, PO Box 829, Sanderton High Wycombe, Bucks, HP14 4HH, UK; 4Institute for Health Services Research and Policy Studies, Northwestern University, 1001 University Place, Evanston, Illinois, IL 60201, USA; 5John Radcliffe Hospital, Headington, Oxford, OX3 9DU, UK; 6Ospedale S. Luigi, Orbassano, Via Regione Gonzole 10, Orbassano, 10126 Italy; 7University of Rostock, Postsch 100888, Rostock, 18055 Germany

Cancer-related anaemia is associated with a wide spectrum of symptoms that can negatively affect quality of life. Because epoetin alfa has demonstrated efficacy in correcting cancer-related anaemia, the impact of this treatment on quality of life was evaluated in a multinational, randomised, double-blind, placebo-controlled trial in 375 anaemic cancer patients receiving non-platinum-based chemotherapy. The cancer-specific measures of quality of life included the general scale (FACT-G Total) and fatigue subscale (FACT-An Fatigue subscale) of the Functional Assessment of Cancer Therapy-Anaemia and the Cancer Linear Analogue Scales measuring energy, ability to do daily activities, and overall quality of life. These measures were also used to examine the relationship between haemoglobin levels and quality of life. Both univariate and multiple linear regression analyses of quality of life data were performed. Results of the univariate analysis have been reported previously. The a priori-planned multiple linear regression analysis, which accounted for the effects of disease progression and several other possibly confounding variables on quality of life, showed a significant advantage for epoetin alfa over placebo for the five scales (all, P < 0.05), and confirmed the results of the univariate analysis. For cancer-specific measures, significant correlations were demonstrated between baseline haemoglobin and quality of life (r, range: 0.14–0.26, all P < 0.05) and between change in haemoglobin and change in quality of life (r, range: 0.26–0.34, all P < 0.01). These findings provide evidence that increasing haemoglobin levels by epoetin alfa administration can significantly improve cancer patients’ quality of life.

British Journal of Cancer (2002) 87, 1341–1353. doi:10.1038/sj.bjc.6600657 www.bjcancer.com © 2002 Cancer Research UK

Keywords: anaemia; cancer; epoetin alfa; haemoglobin; non-platinum chemotherapy; quality of life

Fatigue has replaced pain, nausea, and vomiting as the most common and often the most troubling side effect of concern to cancer patients receiving chemotherapy, with cancer patient surveys commonly and often the most troubling side effect of concern to cancer patients. Moreover, in the studies that evaluated QoL, epoe-
tin alfa provided benefits ranging from improvement in energy and activity levels to an overall improvement in QoL. (Abels, 1992; Littegeb et al., 1994; Glasp'y et al., 1997; Demetri et al., 1998; Gabriole et al., 2001). Epoetin alfa has also been shown to cross the blood–brain barrier and exhibit neuroprotective effects in preclinical studies (Brines et al., 2000; Cerami et al., 2001); and clinical trials in anaemic patients with chronic renal failure suggest that correction of anaemia with epoetin alfa improves brain and cognitive function (Marsh et al., 1991; Pickett et al., 1999).

The results from a study evaluating the effects of epoetin alfa on transfusion requirements and haematopoeitic response, together with a univariate analysis of the study’s QoL data, have been published in detail elsewhere (Littlewood et al., 2001). However, univariate analysis did not take into account possible between-group differences in factors that could influence QoL outcome. Therefore, to assess more precisely the effects of epoetin alfa treatment on QoL, these data were additionally examined by a priori planned multiple linear regression analysis, which controlled for disease progression and other possibly confounding variables (e.g., demographic and baseline clinical characteristics). In this study, we present the findings of the multiple linear regression analysis and explore more fully the relationship between Hb level and QoL. We also take this opportunity to describe the design and results from this study more fully than was possible in the article by Littlewood et al. (2001).

PATIENTS AND METHODS

Patients and study design

The design of the study has been described previously (Littlewood et al., 2001). Briefly, the randomised, double-blind, placebo-controlled trial was conducted in 15 countries at 73 sites with 375 patients (intent-to-treat (ITT) population). The QoL analysis population comprised those ITT patients who had both a baseline and at least 1 follow-up QoL assessment.

To be enrolled, patients were required to have a confirmed diagnosis of solid or non-myeloid haematologic malignancy for which they may or may not have been receiving non-platinum chemotherapy, but were scheduled to receive (an additional) 12 – 24 weeks (3 – 6 cycles) of such therapy. Only patients receiving non-platinum regimens could be included in this phase III study because epoetin alfa was already licensed worldwide for use in patients receiving platinum regimens. In addition, at screening (approximately 1 week prior to randomisation to study drug), patients were required to have Hb levels <10.5 g dL\(^{-1}\), or levels >10.5 g dL\(^{-1}\) but ≤12 g dL\(^{-1}\) following a ≥1.5-g dL\(^{-1}\) decrease in Hb per cycle or month since starting chemotherapy. Patients with acute leukaemia or myeloid malignancies were excluded, as were those with untreated iron, folate, or vitamin B\(_12\) deficiency. Also excluded were patients who had acute major infection or bleeding within 1 month, radiation therapy or allogeneic blood transfusion within 2 weeks, or surgery or acute major illness within 1 week prior to study entry. All patients provided written informed consent before undergoing any study-related procedures.

Patients were stratified by tumour type (solid or haematologic) and Hb level (≤10.5 g dL\(^{-1}\), or >10.5 but ≤12 g dL\(^{-1}\)) at screening. They were randomised in a 2:1 ratio, balanced by using permuted blocks, to receive epoetin alfa 150 – 300 IU kg\(^{-1}\) three times per week or matching placebo by subcutaneous injection. (Outside of the United States, epoetin alfa is manufactured by Ortho Biologics, LLC and distributed and marketed as EPREX\({ }^\text{®}\) or ERYPO\({ }^\text{®}\) by Ortho Biotech and Janssen-Cilag. In the United States, epoetin alfa is manufactured by Amgen Inc. and distributed and marketed by Ortho Biotech Products, L.P.) Study treatment (i.e., epoetin alfa or placebo) was administered for the duration of the study, a maximum of 28 weeks. Study duration included 12 – 24 weeks (3 – 6 cycles) of chemotherapy and a 4-week period when study treatment continued after the last dose of chemotherapy.

The sample size for the trial was determined in consideration of finding a significant effect for the primary efficacy endpoint (transfusions after the first 28 days on-study). Specifically, the sample size calculation was based on detecting an odds ratio of 2 between success and treatment, where success is defined as the absence of any transfusion after the end of the first 4 weeks of treatment. Given a power of 90\%, a significance level of .05 (one-sided), and based on the Cochran-Mantel-Haenszel test, the sample size needed was a priori calculated as 120 in the placebo arm and 240 in the epoetin alfa arm, resulting in a total sample size of 360 (patients randomised to epoetin alfa and placebo in a ratio of 2:1).

Evaluation of quality of life

The QoL hypothesis for this study was that administration of epoetin alfa would improve QoL relative to placebo due to treatment-induced improvements in Hb levels. Clinical efficacy endpoints included the proportion of patients transfused after the first 4 weeks of anaemia treatment (primary endpoint) and change in Hb level from baseline to last available value (secondary endpoint). Change in QoL from baseline to last available assessment was also a secondary efficacy endpoint. The effects of epoetin alfa on QoL were assessed using three patient-rated QoL instruments: subscales of the Functional Assessment of Cancer Therapy-Anaemia (FACT-An), the Cancer Linear Analogue Scale (CLAS, also known as Linear Analogue Scale Assessment, or LASA), and the Medical Outcomes Study Short Form-36 (SF-36). The FACT-An is a 47-item cancer-specific questionnaire comprising a core 27-item general questionnaire (FACT-G Total) that measures physical, social/family, emotional, and functional well-being, and an additional 20-item anaemia questionnaire that assesses the impact of fatigue (13 items, fatigue subscale (FACT-An Fatigue subscale)) and anaemia (seven items, anaemia subscale (FACT-An Anaemia subscale)). The CLAS, another cancer-specific instrument, consists of three QoL visual analogue scales that measure energy level, ability to do daily activities, and overall QoL using a 100-mm scale. The SF-36 is a general health-assessment instrument that provides scores for eight QoL dimensions as well as summary scores for two derived measures (Physical Component Summary and Mental Component Summary (PCS and MCS, respectively)). Possible scores from the FACT-G Total scale range from 0 to 108; for the FACT-An Fatigue subscale, scores range from 0 to 52; and for the three CLAS items scores range from 0 to 100. The SF-36 PCS and MCS scales have no meaningful lower or upper bounds, but are standardised to have a mean of 50 and a standard deviation of 10 points based upon a representative sample from the United States population. The procedure for deriving the standardised PCS and MCS scores has been cross-culturally validated by the instrument developers (Bullinger et al., 1998). Higher scores represent better QoL in every case.

Five QoL scales (FACT-G Total, FACT-An Fatigue subscale (Cella, 1997; Demetri et al., 1998), CLAS: Energy, CLAS: Ability to Do Daily Activities, CLAS: Overall QoL (Cella, 1997; Glasp'y et al., 1997; Demetri et al., 1998)) have demonstrated sensitivity to Hb levels and were therefore designated a priori as the primary QoL efficacy endpoints for analyses of the effects of epoetin alfa. The SF-36 PCS and MCS scales were additionally designated as primary QoL endpoints for analyses, and were included as general measures of QoL.

Patients self-administered the three QoL questionnaires as a single packet a maximum of four times during the study: (1) pretreatment (at the time of randomisation to either epoetin alfa or placebo); (2) before the start of the second on-study chemotherapy cycle for patients on a cyclic chemotherapy regimen (or...
covariance models were estimated for each of the seven primary factors in each model elucidates pure treatment effect of epoetin alfa on QoL scores at follow-up (Demetri et al., 2001). Multiple linear regression analyses were performed as a validation test for previous studies indicating that disease progression has a profound effect on QoL during chemotherapy. However, chemotherapy may have been discontinued earlier as a result of disease progression or toxicity of the chemotherapeutic regimen. Therefore, some patients who did not begin a second cycle of chemotherapy, and thus did not self-administer the first follow-up QoL assessment, may still have completed the epoetin alfa study. In such cases, patients were asked to complete the QoL assessment scheduled for study completion. Because patients could be enrolled with a variable number of expected chemotherapy cycles, and because those cycles could be of varying durations, the timing of QoL assessments could not be standardised to specific days or weeks in the study. The actual timing of the QoL assessments that resulted from this study design is illustrated in Figure 2, where the vertical axis represents patients (sorted by days in the study) and the horizontal axis represents study duration. Missing assessments were not imputed.

Statistical methods

For all analyses, QoL change scores were calculated by subtracting the baseline score from the score obtained at the last available assessment. Changes in QoL data were examined by univariate and multiple linear regression analyses. Univariate analysis methods have been published elsewhere (Littlewood et al., 2001). Multiple linear regression analyses, which were planned a priori, were performed because previous studies indicated that disease progression has a profound effect on QoL at follow-up (Demetri et al., 1998). Therefore, including disease progression and other confounding factors in each model elucidates pure treatment effect on QoL by adjusting for the effects of these factors. These multiple linear regression analyses were performed as a validation test for the originally published univariate analyses.

For the multiple linear regression analyses, separate analyses of covariance models were estimated for each of the seven primary endpoints using the SAS General Linear Model (GLM) procedure (SAS Institute, Inc., Cary, NC, USA). In all models, the QoL change score was the dependent variable. Independent variables included treatment with epoetin alfa, disease progression, treatment interaction with disease progression, and several selected demographic and clinical variables (age, sex, race, baseline Hb level, baseline neutrophil count, baseline reticulocyte count, baseline erythropoietin level, prestudy transfusion dependence, and tumour type (solid or haematologic)). Variables representing treatment with epoetin alfa and disease progression and its interaction with treatment were included in every model. Response to chemotherapy was collected at study termination or early withdrawal. A progressive disease flag was constructed from the chemotherapy response values by setting the flag to one if the response was Progressive Disease (increase in estimated tumour mass by >25% or appearance of new lesion), and by setting the flag to zero otherwise. Patients who were not assessed were categorised as ‘Unknown’ (i.e., not progressive disease). All other independent variables were retained in the model if they were significant at or below the 10% level. Mean QoL change was estimated for each treatment group from the regression parameter estimates using the Least Squares Means (LSM) method (Searle et al., 1980). T statistics were then generated to test for differences in mean change scores between the treatment groups. All significance tests were two-sided, with the alpha level set at 0.05, adjusting for multiple comparisons using a sequentially rejective version of the Bonferroni procedure ( Hochberg, 1988).

The LSM analysis was performed over the entire QoL analysis period adjusted for all covariates, including disease progression. From this analysis, the overall mean changes in QoL for...
the two treatment groups were estimated. However, because disease progression and its interaction with treatment were included in each multiple linear regression model, LSMS of adjusted QoL change scores could also be generalised to two separate patient populations: a population containing only patients exhibiting disease progression and another containing only patients not exhibiting disease progression (i.e., stable, partial, or complete responders to chemotherapy). It should be emphasised that the LSM analysis of patients exhibiting disease progression was not a subgroup analysis; it was an estimate resulting from the regression parameters calculated over all study patients in the entire QoL cohort. The results of such an analysis can be interpreted as predicting the treatment-related QoL benefit that would be expected for a hypothetical patient exhibiting disease progression or for a hypothetical patient not exhibiting disease progression. Of course, in practice, clinicians cannot generally predict which patients will progress. However, because disease progression has such a detrimental effect upon patient QoL, we have explicitly included that possibility in our model specification. This addresses the criticism recently made by Bottomley et al that most QoL papers investigating the effects of epoetin alfa do not take tumour response into account (Bottomley et al, 2002).

Given the underlying QoL hypothesis that administration of epoetin alfa would improve QoL relative to placebo due to treatment-induced improvements in Hb levels, Pearson correlation coefficients were calculated to assess the relationship between baseline Hb level and QoL (cross-sectional correlation), and between change in Hb level and change in score for each primary QoL endpoint (longitudinal correlations). F values associated with these correlation coefficients were also adjusted for multiple comparisons using the sequentially rejective Bonferroni procedure.

RESULTS

Table 1 presents the QoL assessments expected at baseline and follow-up based upon the study design and timing of discontinuing patients. There was a minimal amount of avoidable QoL data missing in this trial due to administrative or logistical reasons (approximately 6.5%, shown as missing CLAS data in Table 1). As seen in the last column of Table 1, 1076 of 1151 QoL assessments expected were available. In the context of clinical studies in oncology, this represents a very low rate of avoidable missing data (Bernhard et al, 1998).

Generally, there was a very low rate of missing QoL items (as opposed to multi-item scales), given the completion of an assessment (Table 2). Over the seven QoL scales chosen a priori for analysis, there was approximately a 2.45% rate of missing items. The FACT-An and the SF-36 scales can be scored in the presence of some level of missing items, but those multi-item scales do mandate a requisite number of non-missing items (usually 50% non-missing). Thus, the missing scale rates for the seven QoL scales chosen a priori for analysis are also presented in Table 2. As the table shows, there was a 2.09% missing scale rate for these seven scales combined. Because the SF-36 PCS and MCS are derived from the same set of items, the percent missing for these two scales is equivalent.

In addition to the data missing due to administrative or logistical reasons, 54 patients in the ITT population had no FACT or SF-36 data owing to unavailability of these questionnaires in some languages (i.e., Czech, Greek, Hungarian, Polish), and an additional 23 were missing those data for other reasons. The CLAS questionnaire was available and administered in the languages named above, as well as in Dutch, English, French, German, Italian, and Portuguese, but 39 patients had missing CLAS data for non-language-related reasons. Therefore, a total of 298 (79.5%) and 336 (89.6%) patients from the ITT population were included in the FACT and CLAS analyses, respectively. The numbers of patients from each country whose FACT or CLAS data were available for analysis are shown in Table 3. Figure 3 is a flow diagram, recommended for use by the CONSORT statement, that displays patients’ progress through the phases of the trial (Moher et al, 2001).

### Table 1 Rates of missing items and scales across QoL assessments

| Scale | Per cent missing items | Per cent missing scales |
|-------|------------------------|-------------------------|
| FACT–G total | 3.87 | 3.52 |
| FACT–An Fatigue subscale | 1.58 | 1.14 |
| CLAS: Energy | 0.00 | 0.00 |
| CLAS: Daily activities | 0.09 | 0.09 |
| CLAS: Overall QoL | 0.18 | 0.18 |
| SF-36 PCS | 1.86 | 5.30 |
| SF-36 MCS | 1.86 | 5.30 |
| Total | 2.45 | 2.09 |

### Table 1 Per cent missing QoL (CLAS) assessments

|                          | Baselinea | First follow-up | Second follow-up | Third follow-upb | Total follow-up |
|--------------------------|-----------|----------------|-----------------|-----------------|----------------|
| QoL assessments expected | 375       | 351            | 294             | 132             | 1152           |
| Died without assessments | 1         | 0              | 0               | 0               | 1              |
| Net assessments expected | 374 (100%)| 351 (100%)    | 294 (100%)      | 132 (100%)      | 1151 (100%)    |
| Missing assessments      | 18 (4.8%) | 29 (8.3%)     | 29 (8.2%)       | 20 (3.0%)       | 128 (6.5%)     |
| Discontinuations         | 16        | 30             | 62              | 20              | 128            |
| Died with assessments    | 6         | 18             | 6               | 0               | 30             |
| Completers              | 1         | 9              | 94              | 112             | 216            |
| Remainder of assessments | 351       | 294            | 132             | 0               | 1076 (93.5%)   |
| Assessments available   | 356 (95.2%)| 322 (91.7%) | 270 (91.8%)     | 128 (97.0%)     | 1076 (93.5%)   |

aBefore first on-study chemotherapy cycle; bWithin ±3 days of start of second on-study chemotherapy cycle; cWithin ±3 days of start of fifth on-study chemotherapy cycle; dAt study completion/withdrawal.
than 12 g dL$^{-1}$ by the time their baseline measurements were assessed. These patients were included in the study and in the ITT analyses.

Transfusion needs and haematopoietic response

Fewer epoetin alfa-treated patients than placebo-treated patients in the ITT population were transfused after day 28 (24.7% vs 39.5%, $P<0.01$, Wald's $\chi^2$ test from logistic model correcting for tumour type and Hb stratum) (Littlewood et al, 2001). (For this analysis, patients on study 28 days or less were counted as transfused, that is, as failures.) Patients treated with epoetin alfa had a significantly greater increase in mean ($\pm$ s.d.) Hb level than did those treated with placebo ($2.2 \pm 2.18$ g dL$^{-1}$ vs $0.5 \pm 1.79$ g dL$^{-1}$, $P<0.01$, $t$-test). (Transfusion-related changes in Hb level were included in the calculation of mean change.) This pattern was observed regardless of tumour type (solid vs haematologic) or Hb stratum (Littlewood et al, 2001).

### Table 3 QoL population by country of enrolment

| Country               | ITT n (%) | FACT-An n (% of ITT) | CLAS n (% of ITT) | FACT-An or CLAS n (% of ITT) |
|-----------------------|-----------|-----------------------|-------------------|------------------------------|
| Germany               | 77 (20.53)| 72 (93.51)            | 72 (93.51)        | 72 (93.51)                   |
| The Netherlands       | 60 (16.00)| 54 (90.00)            | 51 (85.00)        | 54 (90.00)                   |
| Great Britain/Ireland | 42 (11.20)| 40 (95.24)            | 38 (90.48)        | 40 (95.24)                   |
| Belgium/Luxembourg    | 36 (9.60 )| 28 (77.78)            | 25 (69.44)        | 28 (77.78)                   |
| Italy                 | 30 (8.00 )| 30 (100.00)           | 30 (100.00)       | 30 (100.00)                  |
| South Africa          | 25 (6.66 )| 25 (100.00)           | 21 (84.00)        | 25 (100.00)                  |
| France                | 21 (5.60 )| 19 (90.48)            | 18 (85.71)        | 19 (90.48)                   |
| Switzerland           | 16 (4.26 )| 16 (100.00)           | 16 (100.00)       | 16 (100.00)                  |
| Poland                | 15 (4.00 )| 0 (0.00)              | 15 (100.00)       | 15 (100.00)                  |
| Portugal              | 14 (3.73 )| 14 (100.00)           | 14 (100.00)       | 14 (100.00)                  |
| Greece                | 17 (4.53 )| 0 (0.00)              | 14 (82.35)        | 14 (82.35)                   |
| Hungary               | 13 (3.46 )| 0 (0.00)              | 13 (100.00)       | 13 (100.00)                  |
| Czech Republic        | 9 (2.40 ) | 0 (0.00)              | 9 (100.00)        | 9 (100.00)                   |
| TOTAL                 | 375 (100.00)| 298 (79.47)       | 336 (89.60)       | 349 (93.07)                  |

Figure 3 Flow diagram of the patients' progress through the phases of the trial.
Changes in quality of life

The multiple linear regression models and parameter values used to generate LSMs for each of the analyses in this report are presented in Table 5. Table 6 presents LSM estimates for the seven primary QoL endpoints for the overall analysis and by disease progression status. Results of both the univariate and multiple linear regression analyses of QoL change scores for the two treatment groups are displayed in Figures 4, 5, and 6 with the previously published (Littlewood et al, 2001) univariate analyses QoL results included as a reference.

After adjusting for possible differences in demographic and clinical characteristics between the treatment groups, results of the multiple linear regression analyses confirmed the results of the univariate analyses. All within-group mean change scores for the epoetin alfa group were again positive (Figures 4 – 6). Further, mean changes for all seven of these scores were statistically significant (range, \( P < 0.01 \) to \( P < 0.05 \)) (Table 6). In contrast, within-group mean change scores for the placebo group were negative, except for the SF-36 MCS and PCS (Figures 4 – 6). For the overall QoL population, between-group comparisons of the LSM QoL change scores showed statistically significant differences favouring epoetin alfa for all five cancer-specific scales (range, \( P = 0.01 \) to \( P = 0.04 \)) but not for the two SF-36 scales (Figures 4 – 6, Table 6). However, the two SF-36 scales showed no evidence of deterioration in overall physical or mental QoL for patients treated with epoetin alfa.

As estimated by the multiple linear regression model, a hypothetical patient treated with epoetin alfa and exhibiting no disease progression can expect significant benefit as measured by the FACT-An Fatigue subscale and all three CLAS scales. However, a hypothetical patient exhibiting disease progression (in this study, approximately 28% of the QoL population experienced disease progression) could expect no benefit in these measures as based upon the multiple linear regression results (Table 6). These results can be seen as validating the sensitivity of the QoL instruments.

---

Table 4  Baseline characteristics by covariate (QoL population)

| Covariate                  | Epoetin Alfa (n=238) | Placebo (n=111) |
|----------------------------|----------------------|-----------------|
| Sex, n (%)                 |                      |                 |
| Female                      | 158 (66)             | 77 (69)         |
| Male                        | 80 (34)              | 34 (31)         |
| Age (years)                 |                      |                 |
| Mean ± s.d.                 | 58.1 ± 14.2          | 59.2 ± 14.3     |
| Range                       | 18.7–84.9            | 21.1–88.6       |
| Race, n (%)                 |                      |                 |
| White                       | 230 (97)             | 106 (96)        |
| Asian                       | 1 (<1)               | 3 (3)           |
| Black                       | 4 (2)                | 0 (0)           |
| Other                       | 3 (1)                | 2 (2)           |
| Haemoglobin (g dl\(^{-1}\))|                      |                 |
| Mean ± s.d.                 | 9.9 ± 1.1            | 9.8 ± 1.1       |
| Median                      | 10.0                 | 9.7             |
| Range                       | 5.9–14.3             | 6.6–12.7        |
| Tumour type, n (%)          |                      |                 |
| Solid                       | 127 (53)             | 60 (54)         |
| Haematologic                | 111 (47)             | 51 (46)         |
| Neutrophil count, %         |                      |                 |
| Mean ± s.d.                 | 61.5 ± 20.9          | 62.1 ± 20.4     |
| Median                      | 66.0                 | 67.4            |
| Range                       | 1.0–94.5             | 2.0–96.0        |
| Reticulocyte count, %       |                      |                 |
| Mean ± s.d.                 | 2.2 ± 1.3            | 2.4 ± 1.5       |
| Median                      | 2.0                  | 2.1             |
| Range                       | 0.0–7.5              | 0.0–11.5        |
| Serum erythropoietin, mU mL\(^{-1}\)| |                 |
| Mean ± s.d.                 | 102.9 ± 196.8        | 94.1 ± 108.0    |
| Median                      | 49.0                 | 50.0            |
| Range                       | 10–1990              | 10–597          |

Figure 4  QoL mean change scores by treatment group: results of univariate (Littlewood et al, 2001) and multiple linear regression analyses* (FACT-G Total scale and FACT-Fatigue Subscale). *P values adjusted for multiple comparisons (sequentially rejective Bonferroni procedure).
used in this study. One would expect that in a general population of cancer patients (i.e., a population containing both responders and non-responders to cancer treatment), a significant degradation in patient QoL would generally be associated with significant disease progression, regardless of cancer-related anaemia treatment.

Correlation between haemoglobin and quality of life

Correlation analyses were performed to examine the relationship between Hb level and QoL (Table 7). Positive cross-sectional correlations were found for every scale except the SF-36 MCS scale. Slightly stronger longitudinal correlations were found using all seven primary QoL scales; all P values were significant (range, \( P < 0.01 \) to \( P = 0.03 \)). Figure 7 demonstrates the relationship between Hb and QoL over time by showing biweekly Hb levels (Littlewood et al, 2001) and monthly CLAS Energy levels. While the QoL assessments were not designed in this study to yield monthly mean scores and no analysis of monthly mean scores was conducted or reported in this study, this graph does help to visualise the longitudinal relationship between Hb and QoL.
### Table 5 QoL change score multiple linear regression parameter estimates

| Covariates                  | FACT-G Total | FACT-An Fatigue | Energy | Activities | Overall | PCS   | MCS   |
|-----------------------------|--------------|-----------------|--------|------------|---------|-------|-------|
| Sample Size                 | n=253        | n=266           | n=312  | n=284      | n=311   | n=237 | n=225 |
| Placebo                     | n=175        | n=185           | n=215  | n=196      | n=215   | n=160 | n=153 |
| Regression analyses         |              |                 |        |            |         |       |       |
| R²                          | 0.23         | 019             | 0.15   | 0.20       | 0.17    | 0.25  | 0.11  |
| Intercept                   | 6.37         | 7.31            | -3.72  | 72.96b     | -0.84   | -0.32 | 20.79b|
| Solid tumour group          |              |                 |        |            |         |       |       |
| Epoetin alfa                | 4.58b        | 6.64b           | 15.05b | 13.95b     | 14.50b  | 2.92b | 2.03  |
| Disease progression         | -9.57b       | 3.05            | -7.68  | -12.09     | -4.95   | -3.33 | -4.13 |
| Epoetin alfa * disease      | -0.30        | -8.00b          | -15.50 | -14.44     | -20.61b | -5.97b| -1.77 |
| Placebo                     | 5.92b        | 2.68            | 5.78   | 6.47       | 5.67    | 4.10  | 2.76  |
| Placebo                     | 4.34b        | 0.68            | 3.92   | 4.95       | 4.79    | 8.29  | 7.94  |
| CAUcian                     | 12.88b       | 10.66b          | 23.68b | 17.23      | 29.04b  | 9.84b |       |
| Age                         |              |                 |        |            |         |       |       |
| Female                      |              |                 |        |            |         |       |       |
| Baseline haemoglobin        | -1.28        | -1.29b          | -2.05  | -5.64b     | -3.01b  |       |       |
| Baseline neutrophil count   | -0.71        | -0.96           | -1.12  | -2.06      | -0.57a  |       |       |
| Baseline reticulocyte count | -1.18b       |                 |        |            | -0.97a  |       |       |
| Baseline endogenous EPO     |              |                 |        |            |         |       |       |
| Pre-study transfusion       |              |                 |        |            |         |       |       |
| Solid tumour group          | -5.71b       |                 |        |            | -1.76a  |       |       |

s.e=standard error of the mean, 95% confidence interval. aP≤0.05; bP≤0.01.

### Table 6 Regression analyses: least squares means, last assessment change scores

| QoL Scale                  | Epoetin alfa | Placebo | Difference          |
|----------------------------|--------------|---------|---------------------|
|                             | Mean | s.e. | Nom. P value | Adj. P value | Mean | s.e. | Nom. P value | Adj. P value | Mean | s.e. | Nom. P value | Adj. P value |
| Overall                    |      |      |              |              |      |      |              |              |      |      |              |              |
| FACT-G Total               | 2.01 | 0.98 | 0.04         | 0.04         | -2.48 | 1.47 | 0.09         | 0.66         | 4.49 | 1.77 | 0.01         | 0.04         |
| FACT-An Fatigue subscale   |      | 2.70 | 0.84         | <0.01        | -1.70 | 1.27 | 0.18         | 0.90         | 4.40 | 1.52 | <0.01        | 0.02         |
| CLAS: Energy               |      | 7.17 | 1.86         | <0.01        | -3.40 | 2.80 | 0.22         | 0.90         | 10.57 | 3.36 | <0.01        | 0.01         |
| CLAS: Daily Activities     |      | 7.78 | 2.09         | <0.01        | -1.96 | 3.16 | 0.54         | 0.96         | 9.74 | 3.80 | 0.01         | 0.04         |
| CLAS: Overall QoL          |      | 4.46 | 1.82         | 0.02         | -4.10 | 2.76 | 0.14         | 0.83         | 8.56 | 3.31 | 0.01         | 0.04         |
| SF-36: PCS                 |      | 1.27 | 0.60         | <0.01        | 0.05 | 0.87 | 0.96         | 0.96         | 1.22 | 1.06 | 0.02         | 0.33         |
| SF-36: MCS                 |      | 1.88 | 0.87         | <0.01        | 0.35 | 1.28 | 0.78         | 0.96         | 1.53 | 1.55 | 0.33         | 0.33         |
| No disease progression     |      |      |              |              |      |      |              |              |      |      |              |              |
| FACT-G Total               | 4.83 | 1.14 | <0.01        | <0.01        | 0.25 | 1.81 | 0.89         | 0.89         | 4.58 | 2.13 | 0.03         | 0.06         |
| FACT-An Fatigue subscale   |      | 5.79 | 0.97         | <0.01        | -0.85 | 1.54 | 0.58         | 0.89         | 6.64 | 1.82 | <0.01        | <0.01        |
| CLAS: Energy               |      | 13.86 | 2.16   | <0.01       | -1.19 | 3.45 | 0.73         | 0.89         | 15.05 | 4.07 | <0.01        | <0.01        |
| CLAS: Daily Activities     |      | 15.53 | 2.41   | <0.01       | 1.58 | 3.98 | 0.69         | 0.89         | 13.95 | 4.65 | <0.01        | <0.01        |
| CLAS: Overall QoL          |      | 11.83 | 2.11   | <0.01       | -2.67 | 3.41 | 0.43         | 0.89         | 14.50 | 4.01 | <0.01        | <0.01        |
| SF-36: PCS                 |      | 3.92 | 0.69         | <0.01        | 1.00 | 1.47 | 0.35         | 0.89         | 2.92 | 1.27 | 0.02         | 0.06         |
| SF-36: MCS                 |      | 3.55 | 0.99         | <0.01        | 1.52 | 1.60 | 0.34         | 0.89         | 2.03 | 1.88 | 0.28         | 0.28         |
| Disease progression        |      |      |              |              |      |      |              |              |      |      |              |              |
| FACT-G Total               | -5.04 | 1.98 | 0.01         | 0.02         | -9.31 | 2.53 | <0.01        | <0.01        | 4.27 | 3.16 | 0.18         | 0.94         |
| FACT-An Fatigue subscale   |      | -5.26 | 1.64   | <0.01       | -3.90 | 2.20 | 0.08         | 0.21         | -1.36 | 2.74 | 0.62         | 0.94         |
| CLAS: Energy               |      | -9.32 | 3.67   | 0.02        | -8.87 | 4.64 | 0.06         | 0.21         | -0.45 | 5.91 | 0.94         | 0.94         |
| CLAS: Daily Activities     |      | -10.99 | 4.19   | 0.02        | -10.51 | 5.06 | 0.04         | 0.21         | -0.48 | 6.49 | 0.94         | 0.94         |
| CLAS: Overall QoL          |      | -13.73 | 3.59   | 0.01        | -7.62 | 4.54 | 0.09         | 0.21         | -6.11 | 5.79 | 0.29         | 0.94         |
| SF-36: PCS                 |      | -5.38 | 1.20         | <0.01        | -2.33 | 1.47 | 0.11         | 0.21         | -3.05 | 1.89 | 0.11         | 0.75         |
| SF-36: MCS                 |      | -2.36 | 1.82         | 0.20         | -2.61 | 2.09 | 0.21         | 0.21         | 0.25 | 2.71 | 0.93         | 0.94         |

Nom, nominal; Adj, adjusted. aAdjusted using a sequentially rejective Bonferroni procedure; bAs estimated from multiple linear regression across all patients.
The monthly CLAS Energy levels shown in this graph were derived using 28-day windows for months, with the last observation carried forward for missing values and dropouts. It is possible that the decline in QoL seen in the placebo arm, despite relatively stable but low Hb levels, may suggest that long-term anaemia has a cumulative effect on QoL, especially during chemotherapy.

The significant correlations seen in Table 7, and QoL response to Hb shown in Figure 7, suggest that the administration of epoetin alfa in this patient population improves health-related QoL, which is associated with treatment-induced improvements in Hb levels.

Safety

Information regarding adverse events has been reported in detail elsewhere (Littlewood et al., 2001). Epoetin alfa was well tolerated, and the overall incidence of adverse events and the incidence of individual adverse events were generally similar between the two treatment groups.

DISCUSSION

Assessment of QoL has now become a standard, and in some cases mandatory, endpoint in oncology clinical trials and studies. This increased focus on QoL as a therapeutic outcome is being driven by advances made in cancer treatment and management over the last few decades. Earlier detection of malignancies and widespread availability of cancer screening programmes have increased the possibility of successful treatment and recovery, although often at the cost of functional ability and physical, emotional/mental, or social well-being. Also, more intense treatment strategies are being employed with the potential for greater toxicity and associated negative impact on QoL. Determination of the QoL outcome for any potential therapy, whether cancer-specific or supportive, is therefore essential for evaluating comparative treatments as well as making decisions about future treatment or palliative care (Cella, 1997). The primary therapeutic endpoints of concern to the clinical scientist may be very different from those of patients and, consequently, the value placed on gaining relief from symptoms may differ also. Thus, to make informed decisions about the benefits of supportive treatments that have cost implications, both doctors and their patients need the information gained from...
comprehensive QoL evaluation, not merely clinical or economic evaluation.

The FACT-G Total, FACT-An Fatigue subscale, CLAS, and SF-36: PCS and MCS scales were chosen as primary QoL endpoints to assess the effects of epoetin alfa on QoL in the present study. The FACT-G Total, FACT-An Fatigue subscale, and CLAS were considered especially appropriate for this purpose as they are cancer-specific, Hb-sensitive, and easily completed by patients. The CLAS, in addition, had been used frequently in previous trials of epoetin alfa and, at the time this study was initiated, was more widely available than the FACT scales in the languages of the participating countries (10 vs six languages). Also, its use could facilitate comparisons between the results of this study and previous trials. The SF-36 is not cancer-specific but was included as a general measure to ensure that there were no unexpected adverse consequences of epoetin alfa therapy affecting overall QoL.

To clarify the effect of epoetin alfa on QoL, beyond that originally reported based upon univariate analyses (Littlewood et al., 2001), a multiple linear regression analysis was performed that took into account tumour progression as well as many other possibly confounding variables (e.g., age, sex, baseline Hb). As in the univariate analysis, the P values were adjusted to account for multiple comparisons. The results of this analysis confirmed the advantage of epoetin alfa over placebo, as demonstrated in the univariate analysis, with significant differences again noted for all cancer-specific scales. While the generality of the two SF-36 summary scales reduced their ability to discriminate between groups in this analysis, the positive (though non-significant) SF-36 changes favouring epoetin alfa strongly suggest that there were no general trends in unforeseen detrimental effects on QoL from epoetin alfa treatment using an instrument that casts a wide net for functional and well-being domains. In fact, it should be noted that the within-group analysis shows a positive and statistically significant change in the two SF-36 scales for the epoetin alfa group, but no corresponding significant change for the placebo group.

While only 2–3% of the patients included in the current report were non-Caucasian, race was included as a covariate because it has historically been a strong predictor of QoL and is customarily included in multiple linear regression analyses of treatment effect on QoL. The variable’s highly significant parameter estimates, despite the low number of non-Caucasian patients included in the study, indicates that race may be an important predictor of patient QoL. For this reason, and to maintain consistency with the main body of QoL research, we decided to retain race in our regression models.

A possible limitation of this analysis is the focus upon the QoL change from baseline to the last available assessment. Analyses incorporating change scores derived in this manner may be affected by bias across treatment arms. For example, patients in one treatment arm may stop early due to anaemia (and thus a low QoL) by bias across treatment arms. For example, patients in one treatment arm may stop early due to anaemia (and thus a low QoL) despite the low number of non-Caucasian patients included in multiple linear regression analyses of treatment effect on cancer-specific QoL. This corroborated by several randomised placebo-controlled and open-label studies (Abels, 1992; Glaspy et al., 1997; Demetri et al., 1998; Gabrilove et al., 2001). In the earliest of these studies (Abels, 1992), QoL was assessed in a series of placebo-controlled trials conducted in three different patient populations: patients receiving no chemotherapy, patients receiving non-cisplatin-containing chemotherapy, and patients receiving cisplatin-containing chemotherapy. QoL was evaluated by means of visual analogue scales that measured energy level, ability to do daily activities, and overall QoL (scales equivalent to the CLAS scales used in the present study). Comparison of QoL scores for the total epoetin alfa population of the three trials with those for the total placebo population revealed significantly (P<0.05) greater improvement in all three QoL parameters in patients who responded to epoetin alfa (i.e., had a haematocrit increase ≥6%) vs patients given placebo. From this, the investigators concluded that epoetin alfa does improve functional capacity in anemic cancer patients when there is an increase in haematocrit.

In the three separate large, open-label, non-placebo-controlled, community-based surveillance studies with a combined enrolment of over 7000 patients, large positive changes were noted in Hb levels and QoL of patients treated with epoetin alfa (Glaspy et al., 1997; Demetri et al., 1998; Gabrilove et al., 2001). Transfusion requirements also decreased steadily and dramatically throughout the course of all three studies. These changes were noted even after accounting for disease progression in two of the studies (Glaspy et al., 1997; Demetri et al., 1998). The relationship between disease progression and QoL changes in patients with anaemia experienced deterioration in QoL. Finally, there were significant correlations noted in these studies between Hb change and a between-group difference of 8.5–10.6 points for the three CLAS scales. The question of the clinical relevance of these changes can be partially answered by the results of two recent analyses. An analysis based on the Littlewood et al study compared the FACT-An and CLAS score changes of improved and stable patients; differences in corresponding QoL scores were considered to be the minimally important differences (MIDs) (Patrick et al., 2000). For the FACT-G Total and FACT-An Fatigue subscale, the MIDs were 2.25 and 4.23, respectively. For the CLAS scales, the MIDs ranged from 8.98 to 10.26. The between-group differences observed from the multivariate analyses exceeded the MIDs for all scales except the CLAS Overall QoL. A separate analysis compared the FACT-An scores from the Littlewood et al (2001) trial with scores from a demographically representative United States sample (Nortier et al., 2001). The analysis found that the between-group QoL advantage seen in epoetin alfa-treated patients represented 95% and 51% of the deficit from normative scores seen at baseline for the FACT-G Total and FACT-An Fatigue subscale, respectively. Based on the conclusions of these analyses, the results for the cancer-specific scales seen in the clinical trial appear not only to be statistically significant, but also clinically relevant.

Due to space limitations and, often, trial design, QoL results (if they are published at all) are usually presented in a fashion similar to that used for physiologic endpoints (i.e., a simple statistical comparison of ITT groups). The current study is given to rigorous analysis including multiple linear regression on over a dozen covariates, consideration of patterns of missing data, and further interpretation of the robustness of the results based upon these analyses. Further, the finding of a positive within-group epoetin alfa treatment effect on cancer-specific QoL is corroborated by several randomisation placebo-controlled and open-label studies (Abels, 1992; Glaspy et al., 1997; Demetri et al., 1998; Gabrilove et al., 2001). In the earliest of these studies (Abels, 1992), QoL was assessed in a series of placebo-controlled trials conducted in three different patient populations: patients receiving no chemotherapy, patients receiving non-cisplatin-containing chemotherapy, and patients receiving cisplatin-containing chemotherapy. QoL was evaluated by visual analogue scales that measured energy level, ability to do daily activities, and overall QoL (scales equivalent to the CLAS scales used in the present study). Comparison of QoL scores for the total epoetin alfa population of the three trials with those for the total placebo population revealed significantly (P<0.05) greater improvement in all three QoL parameters in patients who responded to epoetin alfa (i.e., had a haematocrit increase ≥6%) vs patients given placebo. From this, the investigators concluded that epoetin alfa does improve functional capacity in anaemic cancer patients when there is an increase in haematocrit.
and change in QoL (Glaspy et al, 1997; Demetri et al, 1998; Gabri- love et al, 2001), even after accounting for the effects of disease progression (Glaspy et al, 1997; Demetri et al, 1998).

Although both the Abels study and the community-based studies support the main finding of the present study, each study had limitations, the most obvious being that the community-based studies were not placebo-controlled. It is, however, an important finding that several large, community-based studies that more closely represent the actual clinical environment corroborate the results seen in the present randomised, placebo-controlled study. The Abels study used visual analogue scales to measure changes in symptoms and QoL, a method that is recognised in pain assessment and other symptom assessment and has sensitivity in cancer-symptom assessment. Since then, new instruments such as the FACT have been developed to augment measurements such as linear analogue scales; and studies using both measures now report positive changes in both instruments with anaemia correction (Glaspy et al, 1997; Demetri et al, 1998). In addition, the Abels study reported differences in results between patients who received placebo and patients who responded to epoetin alfa treatment rather than reporting results in all patients who received epoetin alfa.

The finding in the present study of an important role for Hb levels in increasing and maintaining QoL also is in agreement with the findings of two other studies that investigated the impact of anaemia and fatigue on cancer patients (Cella, 1997; Langer et al, 1997). In one study (Cella, 1997), assessment of QoL based on the FACT-An showed that patients with Hb levels >12 g dL$^{-1}$ had significantly less fatigue, fewer non-fatigue anaemia symptoms, better physical and functional well-being, and higher overall QoL than those with an Hb level $\leq 12$ g dL$^{-1}$ ($P<0.01$ for all). The second study (Langer et al, 1997), using an index based on FACT-Lung subscales, showed a significant correlation ($r=0.38$, $P<0.002$), independent of tumour response, between worsening anaemia and diminishing QoL in patients with non-small-cell lung cancer by their fourth cycle of chemotherapy. Together with data presented here, these studies provide objective evidence that increasing Hb levels in anaemic cancer patients can positively influence their QoL. (Cella, 1997; Glaspy et al, 1997; Langer et al, 1997; Demetri et al, 1998; Gabri-love et al, 2001).

CONCLUSIONS

In summary, the results of univariate analysis of change scores for cancer-specific scales that evaluated QoL in this randomised study in anaemic patients receiving non-platinum-based chemotherapy were confirmed by multiple linear regression analysis, which adjusted for possible differences in demographic and clinical characteristics between the treatment groups as well as tumour response. Correlation analysis revealed a significant positive relationship between change in Hb level and change in all primary cancer-specific QoL endpoints evaluated. These findings indicate that epoetin alfa can improve QoL in anaemic cancer patients undergoing chemotherapy, and this change is associated with increasing Hb levels. Moreover, the results demonstrate that epoetin alfa treatment and disease progression are independent contributors to changes in QoL.

In the increasingly cost-contained and cost-conscious clinical environments in which many European clinicians work, there might well be reluctance to prescribe a relatively expensive product. The importance of successfully treating chemotherapy-induced anaemia by raising Hb levels with epoetin alfa and the concomitant benefits that this brings for patients’ functioning and well-being are now clear. Providers are therefore faced with a situation similar to that experienced with the advent of the 5HT3 antagonists. Although nausea and vomiting had a severely deleterious effect on the quality of patients’ lives and these symptoms were always ranked at the top of lists charting patients’ concerns, some clinicians were either reluctant or unable to prescribe the most efficacious treatments on the grounds of cost. This was eventually seen as a weak argument, as uncontrolled symptoms that detract from patients’ QoL can jeopardise completion of treatment and may have many hidden or unmeasured costs. Consequently, 5HT3 antagonists are now regarded as a standard part of the management of patients undergoing chemotherapy and this has transformed the lives of many patients, making chemotherapy much more tolerable.

Now that nausea and vomiting in chemotherapy have been significantly alleviated, fatigue due to the chemotherapy-induced anaemia has emerged as a major and distressing problem for patients (Curt et al, 2000). It is usually poorly understood, often managed inadequately, and its impact on patients is generally underestimated. We hope that this study will help to raise awareness that epoetin alfa significantly improves Hb levels and reduces the fatigue that so impairs the QoL of anaemic cancer patients.

ACKNOWLEDGEMENTS

The authors would like to thank the original British Journal of Cancer peer reviewers, whose excellent suggestions greatly improved the quality of the manuscript.

REFERENCES

Abels R (1992) Recombinant human erythropoietin in the treatment of the anaemia of cancer. Acta Haematologica 87 (Suppl 1): 4–11
Barrett-Lee PJ, Bailey NP, O’Brien ME, Wager E (2000) Large-scale UK audit of blood transfusion requirements and anaemia in patients receiving cytotoxic chemotherapy. Br J Cancer 82: 93–97
Bernhard J, Cella DF, Coates AS, Fallowfield L, Ganz PA, Moinpour CM, Mosconi P, Osoba D, Simes J, Hurny C (1998) Missing quality of life data in cancer clinical trials: serious problems and challenges. Stat Med 17: 517–532
Bottomley A, Thomas R, van SK, Flechters H, Djulbegovic B (2002) Human recombinant erythropoietin and quality of life: a wonder drug or something to wonder about? Lancet Oncol 3: 145–153
Brines ML, Ghezzi P, Keenan S, Agnello D, de Lanerolle NC, Cerami C, Itri LM, Cerami A (2000) Erythropoietin crosses the blood-brain barrier to protect against experimental brain injury. Proc Natl Acad Sci USA 97: 10526–10531
Glaspy J, Langer A, Demetri G, Cella D, Davis K, Breitbart W, Curt G for the Fatigue Coalition (2001) Cancer-related fatigue: prevalence of proposed diagnostic criteria in a United States sample of cancer survivors. J Clin Oncol 19: 3385–3391
Cerami A, Brines ML, Ghezzi P, Cerami C (2001) Effects of epoetin alfa on the central nervous system. Semin Oncol 28 (2 Suppl 8): 66–70

Bullinger M, Alonso J, Apolone G, Leplege A, Sullivan M, Wood-Dauphinee S, Gandek B, Wagner A, Aaronson N, Bech P, Fukuhara S, Kaasa S, Ware Jr JE (1998) Translating health status questionnaires and evaluating their quality: the IQOLA project approach. J Clin Epidemiol 51: 913–923
Cella D (1997) The Functional Assessment of Cancer Therapy-Anemia (FACT-An) Scale: a new tool for the assessment of outcomes in cancer anaemia and fatigue. Semin Hematol 34 (Suppl 2): 13–19
Cella D (1998) Factors influencing quality of life in cancer patients: anaemia and fatigue. Semin Oncol 25 (Suppl 7): 43–46
Cella D, Davis K, Breitbart W, Curt G for the Fatigue Coalition (2001) Cancer-related fatigue: prevalence of proposed diagnostic criteria in a United States sample of cancer survivors. J Clin Oncol 19: 3385–3391
Cerami A, Brines ML, Ghezzi P, Cerami C (2001) Effects of epoetin alfa on the central nervous system. Semin Oncol 28 (2 Suppl 8): 66–70

© 2002 Cancer Research UK

British Journal of Cancer (2002) 87(12), 1341 – 1353
Coiffier B, Guastalla JP, Pujade-Lauraine E, Bastit P for the Anemia Study Group (2001) Predicting cancer-associated anaemia in patients receiving non-platinum chemotherapy: results of a retrospective survey. *Eur J Cancer* **37**: 1617–1623

Curt GA, Breitbart W, Cell a D, Groopman JE, Horning SJ, Itri LM, Johnson DH, Miaskowski C, Scherr SL, Portenoy RK, Vogelzang NJ (2000) Impact of cancer-related fatigue on the lives of patients: new findings from the Fatigue Coalition. *Oncologist* **5**: 353–360

Demetri GD, Kris M, Wade J, Degos L, Cell a D for the Procrit Study Group (1998) Quality-of-life benefit in chemotherapy patients treated with epoetin alfa is independent of disease response or tumour type: results from a prospective community oncology study. *J Clin Oncol* **16**: 3412–3425

Fairclough D, Gagnon D (2000) The significance of missing data in clinical trials. *ESMOQ: Final Program* **28**: (Abstract 1697)

Gabrilove JL, Einhorn LH, Livingston RB, Sarokhan B, Winer E, Einhorn LH (1998) Fatigue in cancer and HIV/AIDS. *Oncology (Huntingt)* **12**: 335–344

Hochberg Y (1988) A sharpar Bonferroni procedure for multiple tests of significance. *Biometrika* **75**: 800–802

Langer C, Barsewick A, Bruner D, Grindel C, Leighton J, Luckscheiter C, Bonjo C, McAler C (1997) Correlation of quality of life (QoL) with survival, treatment response, and anemia in patients with advanced non-small cell lung cancer (NSCLC) treated with carboplatin and paclitaxel. *Lung Cancer* **18**: 23 (Abstract 79)

Leitgeb C, Pecherstorfer M, Fritz E, Ludwig H (1994) Quality of life in chronic anemia of cancer during treatment with recombinant human erythropoetin. *Cancer* **73**: 2535–2542

Littlewood TJ, Bajetta E, Nortier JW, Vercaemmen E, Rapoport B (2001) Effects of epoetin alfa on hematologic parameters and quality of life in cancer patients receiving nonplatinum chemotherapy: results of a randomized, double-blind, placebo-controlled trial. *J Clin Oncol* **19**: 2865–2874

Ludwig H, Fritz E (1998) Anemia in cancer patients. *Semin Oncol* **25** (Suppl 7): 2–6

Marsh JT, Brown WS, Wolcott D, Carr CR, Harper R, Schweitzer SV, Nissenson AR (1991) rHuEPO treatment improves brain and cognitive function of anemic dialysis patients. *Kidney Int* **39**: 155–163

Moher D, Schulze KF, Altman DG (2001) The CONSORT statement: revised recommendations for improving the quality of reports of parallel-group randomised trials. *Lancet* **357**: 1191–1194

Nortier JWR, Zagarì M, Vandonor, Gagnon D, Cell a D (2001) Epoetin alfa overcomes much of the QOL deficit seen in anemic cancer patients. *Eur J Cancer* **37** (Suppl 6): S531

Patrick DL, Zagarì M, Gagnon D, Rapoport BL (2000) Clinical significance of changes in health-related quality of life (HRQOL) of anemic cancer patients. *Ann Oncol* **11** (Suppl 4): 149

Pickett JL, Theberge DC, Brown WS, Schweitzer SV, Nissenson AR (1999) Normalizing hematocrit in dialysis patients improves brain function. *Am J Kidney Dis* **33**: 1122–1130

Schwartz AI (1998) The Schwartz Cancer Fatigue Scale: testing reliability and validity. *Oncology Nursing Forum* **25**: 711–717

Searle SR, Speed FM, Milliken GA (1980) Populations marginal means in the linear model: an alternative to least squares means. *American Statistical Association* **34**: 216–221

Simon AM, Zitoun R (1999) Fatigue in cancer patients. *Cancer Oncol* **11**: 244–249

Stone P, Richards M, Hardy J (1998) Fatigue in patients with cancer. *Eur J Cancer* **34**: 1649–1651

Stone P, Richardson A, Ream E, Smith AG, Kerr DJ, Kearny N (2000) Cancer-related fatigue: inevitable, unimportant, and untreatable? Results of a multi-centre patient survey. *Cancer Fatigue Forum. Ann Oncol* **11**: 971–975

Vogelzang NJ, Breitbart W, Cell a D, Groopman JE, Horning SJ, Itri LM, Johnson DH, Scherr SL, Portenoy RK (1997) Patient, caregiver, and oncologist perception of cancer-related fatigue: results of a tripart assessment survey. *Int J Radiat Oncol Biol Phys* **34**: 4–12

Winnimming ML, Nail LM, Burke MB, Brophy L, Cimprich B, Jones LS, Pickard-Holley S, Rhodes V, St Pierre B, Beck S et al (1994) Fatigue and the cancer experience: the state of the knowledge. *Oncology Nursing Forum* **21**: 23–36

L Fallowfield

Di Ematologia, Milan; and Gabriella Gorzegno, MD, Osperdale S. Studio E La Cura dei Tumori, Milan; Agostino Cortelezzi, MD, Istituto di Clinica Medica Generale, Orbassano (Torino).

Di Ematologia, Milan; and Gabriella Gorzegno, MD, Osperdale S. Studio E La Cura dei Tumori, Milan; Agostino Cortelezzi, MD, Istituto di Clinica Medica Generale, Orbassano (Torino).
tenberg, Amersfoort; M.R. Schaafsma, MD, Medisch Spectrum Twente, Enschede; L.H. Siegenbeek van Heukclom, MD, Medisch Centrum Alkmaar, Alkmaar; C. Van der Heul, MD, Sint Elisabeth Ziekenhuis, Tilburg; Marinus Van Marwijk Kooy, MD, St Sophia Ziekenhuis, Zwolle; Gerard Vreugdenhil, MD, Sint Joseph Ziekenhuis, Veldhoven; and Jacques Wils, MD, Laurentius Hospital, Roermond. Poland: Jerzy Holowiecki, MD, Department of Haematology, Silesian School of Medicine, Katowice; Marek Pawlicki, MD, Maria Sklodowska-Curie Memorial Cancer Center, Division in Cracow, Kraków; and Piotr Siedlecki, MD, Maria Sklodowska-Curie Memorial Cancer Center, Warsaw, Warszawa. Portugal: Cândida Azevedo, MD, Instituto Português de Oncologia, Porto; Ricardo Marques da Costa, MD, Hospital Dristrital de Leiria, Leiria; and Joaquim Gouveia, MD, Hospital de Santo António dos Capuchos, Lisboa. South Africa: Dayle Hacking, MD, Durban Oncology Center, Westridge, Durban; Johann Raats, MD, 110 Delmar Medical Center, Capetown; and Bernardo Rapoport, MD, The Medical Oncology Center of Rosebank, Johannesburg. Switzerland: Matti S. Aapro, MD, Centre Anticancereux, Genolier; Richard Herrmann, MD, Kantonsprital Basel, Basel; J.M. Lüthi, MD, Regionalsprital Thun, Thun; and Kaspar Rhyner, MD, Kantonsprital Glarus, Glarus. United Kingdom: Peter Jeffrey Barrett-Lee, MD, Department of Oncology, Velindre Hospital, Cardiff; David Fairlamb, MD, New Cross Hospital, Cansly Center, Wolverhampton; Riaz Jan-Mohamed, MD, Hillingdon Hospital, Uxbridge; Timothy James Littlewood, MD, John Radcliffe Hospital, Oxford; Graham Smith, MD, Royal United Hospital, Bath; and John Sweetenham, MD, Southampton General Hospital, CRC Oncology Unit, Southampton.