The development of targeted agents holds considerable promise for cancer treatment but progress to date has not been easy. Many of the difficulties encountered in the development of targeted anticancer agents can be explained by an incomplete understanding of human tumour biology, limited understanding of the drug target and problems with patient selection. The development of targeted anticancer agents has required novel trial designs as well as the investigation of new pharmacodynamic and surrogate trial end points. The future success of targeted therapy will involve initiatives to identify the most promising targets, closer integration of preclinical and clinical data, a greater use of genomic and proteomic techniques, and further refinement of trial design.

SELECTING TARGETED AGENTS FOR TARGETED PATIENT POPULATIONS

Most cancers are driven by and dependent upon multiple aberrant signalling pathways, and thus ‘single-hit’ therapy may not represent an optimal approach in many clinical situations. It is evident that there is scope for significant redundancy in cell signalling and selecting a single target within a heterogeneous tumour type may yield clinically disappointing results. In tumours with a relatively narrow range of critical genetic defects (e.g., acute promyelocytic leukaemia and chronic phase chronic myeloid leukaemia (CML)), the ability to develop successful targeted agents with striking activity has been more easily accomplished than in more complex and heterogeneous tumour types (e.g., breast cancer, and non-small-cell lung cancer (NSCLC)). Most clinically apparent tumours contain multiple genetic defects with incompletely defined phenotypic consequences. The declining therapeutic impact of imatinib as one proceeds in Philadelphia-positive CML (Cheson et al., 2002), illustrates the impact of increasing genetic alterations in conferring drug resistance to a targeted agent (Gorre et al., 2001; von Bubnoff et al., 2002). For successful targeted drug development we ideally need to be able to identify targets that provide a critical transforming signal to the tumour as distinct from targets where expression is not linked with signal-dependency. We must also acknowledge that targeted drugs may have the most marked effects on only a subset of tumour cells and may be incapable of inhibiting quiescent and perhaps non-target-addicted tumour stem cell populations. Targeted drugs may therefore be able to provide tumour control rather than cure.

The clinical evaluation of targeted agents has provided some refinement in the classification of complex tumours, and this trend is likely to gather pace as tumours become defined by molecular features rather than according to light microscopic classification. The emergence of trastuzumab as a useful agent in a subset of breast cancer patients is one example of the impact of targeted treatment in refining tumour classification (Vogel et al., 2002). Likewise, the finding that a small subset of NSCLC patients who have gain-of-function epidermal growth factor receptor (EGFR) mutations and response rates to gefitinib and erlotinib of over 80% must be viewed as a major advance in our understanding of lung cancer biology (Lynch et al., 2004; Paz et al., 2004; Mitsudomi et al., 2005; Tokumo et al., 2005). The future clinical evaluation of targeted agents will shed new light on tumour pathogenesis, provide better understanding of tumour biology and result in changes to the way that tumours are classified.

For trastuzumab (Herceptin®) and imatinib (Glivec®), a limited number of molecular features can be used to define a suitable target patient population, but it seems unlikely that all patients who could derive clinical benefit from targeted drugs will be identified by determination of a single molecular feature. We know also that tumour architecture is heterogeneous and may significantly affect the intratumoural pharmacokinetics of a drug (Jayson et al., 2002). Thus, a drug that is effective at the molecular level may not necessarily achieve clinically useful doses in the tumour.
Complex tumour types may be more susceptible to the use of multiple-targeted agents or to drugs that target generic, fundamental tumour processes such as angiogenesis, apoptosis and cellular proliferation. For example, small molecule tyrosine kinase inhibitors (TKIs) such as ZD6474 (inhibitor of vascular endothelial growth factor receptor (VEGFR) and EGFR) (Wedge et al, 2002; Ciardiello et al, 2003), SU11248 (inhibitor of VEGFR, platelet-derived growth factor receptor (PDGFR), c-Kit, colony-stimulating factor receptor-1 (CSFR-1) and FLT3-R) and vatalanib (PTK787/ZK222584; inhibitor of VEGFR and PDGFR) (Abrams et al, 2003; Thomas et al, 2003) show activity against more than one receptor and are inhibitory in a wide range of preclinical models.

Since there are no predefined predictive rules, a major future challenge must be to identify critical molecular events that drive tumour growth, survival and metastasis. Important determinants of tumour behaviour are likely to be DNA mutational or rearrangement events, which produce oncogenes with dominant gain of function or tumour suppressor genes with recessive loss of function.

**CLINICAL TRIAL DESIGN**

The traditional phase I trial design used for cytotoxic drugs has been recognised as having limitations for targeted agents. Instead of the maximum tolerated dose, definition of the optimum biological dose became a central theme in phase I studies (Gelmon et al, 1999; Korn et al, 2001). However, a retrospective review of 60 phase I trials of targeted agents reported mainly between 1999 and 2002 revealed that the majority of studies had used toxicity as the sole determinant for dose selection in phase II studies. Nontraditional end points such as measurements of drug effect or functional imaging have not been routinely incorporated into phase I trial design and have rarely formed the basis for future dose selection or trial design (Parulekhar and Eisenhauer, 2004).

Proof-of-principle studies to measure drug effect on target are highly desirable during early drug development, but are difficult to perform. Although tumour represents the tissue of primary interest, tumour sampling can be obtained in only a limited proportion of patients and at a small number of time points. Trials with mandatory biopsies have greater ethical and administrative complexity, and the net result is often slower patient recruitment and recruitment of a narrow range of tumour types. Preclinical models are needed that define an appropriate degree of target inhibition for clinical studies, but observations of target inhibition still need to be clinically validated. Surrogate tissues (peripheral blood mononuclear cells, skin, hair follicles) may be useful in that they can be more readily collected than tumour biopsies, but the pharmacodynamic relationship between surrogate and tumour cells needs to be understood and defined. The incorporation of surrogate and tumour pharmacodynamic markers into clinical trials has also highlighted the fact that assay validation, reliability and quality assurance need attention if results are to be interpretable (Colburn, 2003; Cummings et al, 2004).

Once an agent has been found to have acceptable toxicity in phase I (ideally with proof-of-principle that it has achieved inhibition of the putative biological target), it is necessary to demonstrate that it has clinical activity in phase II and III trials. Patient selection for such trials is often problematic, particularly if there is no established link between the chosen target and a ‘diagnostic signature’. Translational research has established a significant correlation between genomic amplification and high expression of HER2/neu in advanced breast cancer, and response and clinical benefit from trastuzumab (Vogel et al, 2002). However, the predictive capacity of this observation has its limitations, since despite the development of validated molecular diagnostics for HER2, only a minority of patients with HER2/neu-positive tumours will actually respond to trastuzumab monotherapy.

However, the relatively high response rates in phase II/III trials of imatinib in metastatic gastrointestinal stromal tumours (GIST) support the intrinsic power of a science-driven approach to patient selection (van Oosterom et al, 2001; Demetri et al, 2002). Imatinib is an inhibitor of c-Kit receptor tyrosine kinase, and constitutive activation of this receptor, resulting from gain-of-function mutations in the c-Kit gene, had been hypothesised to be critical in GIST (Rubin et al, 2001). The situation for EGFR TKIs has however been more challenging. In this case, the target receptor was found to be widely expressed in epithelial tumours and strong preclinical data supported a role for EGFR in epithelial tumour biology. Nevertheless, the relationship between EGFR expression, target inhibition and tumour dependency on EGFR was unclear, and it was clinical observations that propelled a breakthrough in basic science. Responses to gefitinib in advanced NSCLC were encountered in a minority of patients who more frequently had adenocarcinoma, were non- or ex-smokers, and were more likely to be female or Japanese (Fukuoka et al, 2003; Kaneda et al, 2004; Miller et al, 2004). Molecular characterisation of responding patients identified a novel subset of NSCLC patients with gain-of-function mutations in the ATP-binding domain of the receptor (Lynch et al, 2004; Paez et al, 2004; Mitsudomi et al, 2005; Tokumo et al, 2005). Had this biology been known in advance, patient selection based upon this criterion could have yielded a response rate to gefitinib and erlotinib of over 80% in phase II trials in this subset of NSCLC. Analysis of randomised clinical data with erlotinib suggests that this select cohort of mutation-positive patients and not those with wild-type receptor (CSFR-1) and FLT3-R) and vatalanib (PTK787/ZK222584; inhibitor of VEGFR and PDGFR) (Abrams et al, 2003; Thomas et al, 2003) show activity against more than one receptor and are inhibitory in a wide range of preclinical models.

Since there are no predefined predictive rules, a major future challenge must be to identify critical molecular events that drive tumour growth, survival and metastasis. Important determinants of tumour behaviour are likely to be DNA mutational or rearrangement events, which produce oncogenes with dominant gain of function or tumour suppressor genes with recessive loss of function.

**IDENTIFYING AND VALIDATING MARKERS OF TARGET INHIBITION**

Although most phase I trials of targeted drugs incorporate translational science, the same cannot be said of many phase II or III trials despite the fact that this arena is likely to be one of the most fruitful for translational science. The ability to perform analysis at DNA, RNA and protein levels and to correlate this with outcome in phase II and III trials should facilitate our ability to...
reclassify tumours and identify subpopulations of patients who gain from targeted therapeutics. In practice, most patients are willing to have additional investigative studies performed on their tumours in the hope that it may yield important information. It is important that all stakeholders involved in the clinical testing of new anticancer agents understand that the correlation of such data with clinical outcome may be critical for the optimal development and use of current and future targeted therapies. Thus, incorporating translational research into phase II and III trials needs to be recognised as an important future objective.

Dose delivery with cytotoxic drugs is normally heavily influenced by normal tissue toxicity (e.g., myelosuppression and mucositis). It seems a reasonable premise that dosing for targeted therapies should be driven by target effect rather than toxicity. While this does not remove the need to evaluate traditional clinical end points (toxicity, tumour response, time to progression and survival), there is increasing interest in identifying biomarkers of drug effect. With the relatively low toxicity of many targeted agents compared with conventional treatment strategies, the short-term administration of a targeted agent before elective surgery may be a useful method for identifying biomarkers that merit further clinical investigation.

CONCLUSION

There are unprece...ened opportunities to translate our understanding of tumour biology into more significant therapeutic gains. However, we must accept that our understanding of clinical human cancer is incomplete and that there are many obstacles to overcome. Identifying a molecular target that can be inhibited and then simply testing a promising lead compound in unselected patients may not be an effective strategy without determining target effect or conducting adequate translational research. As the numbers of patients treated with targeted agents increases, it will become possible to determine whether significant tumour dependency exists for a specific biological target, at least in a subset of patients. Indeed, phase III trials of bevacizumab have recently validated VEGF as a target in both CRC (Hurwitz et al., 2004). We should attempt to elucidate potential mechanisms whereby tumours circumvent this target dependency, as this may help us in patient selection and finding effective combination therapies. Initial clinical development should focus on proof-of-principle, a demonstration that the drug produces inhibition of the biological target at doses that are well tolerated, and that the consequences of target inhibition can be identified and measured using validated surrogates for clinical benefit. This will require the integration of pharmacokinetics, pharmacodynamics and possibly functional imaging into trials. Selection of a specific population of patients is desirable if adequate clinical or preclinical data exists.

In the absence of data allowing patient enrichment, further preclinical work may be required or significant investment in gathering tumour and correlative material should be undertaken in phase I and II studies. Technologies such as genomics, proteomics and biomarker development represent promising approaches with which to probe for predictive markers. Everyone involved in developing novel targeted anticancer therapies should recognise that the most significant future advances are likely to emerge from the incorporation of such technologies into clinical trials. In short, the future success of targeted therapies will need a coordinated, multidisciplinary team approach.

REFERENCES

Abrams TJ, Murray LJ, Pesenti E, Holway VW, Colombo T, Lee LB, Cherrington JM, Pryer NK (2003) Preclinical evaluation of the tyrosine kinase inhibitor SU11248 as a single agent and in combination with ‘standard of care’ therapeutic agents for the treatment of breast cancer. Mol Cancer Ther 2: 1011 – 1021

Carr KM, Rosenblatt K, Petricoin EF, Liotta LA (2004) Genomic and proteomic approaches for studying human cancer: prospects for true patient-tailored therapy. Hum Genomics 1: 134 – 140

Hurwitz H, Fehrenbacher L, Novotny W, Cartwright T, Hainsworth J, Heim W, Berlin J, Baron A, Griffling S, Holmgren E, Ferrara N, Fye G, Rogers B, Ross R, Kabbavinavar F (2004) Bevacizumab plus irinotecan, fluorouracil, and leucovorin for metastatic colorectal cancer. N Engl J Med 350: 2335 – 2342

Jayson GC, Zweit J, Jackson A, Mulatero C, Julyan P, Ranson M, Broughton L, Wagstaff J, Hakannson L, Groenewegen G, Bailey J, Smith N, Hastings D, Lawrence J, Haroon H, Ward T, McGowan AT, Tang M, Levitt D, Marreaud S, Lehmann FF, Herold M, Zwierzina H (2002) Molecular imaging and biological evaluation of HuMV833 anti-VEGF antibody: implications for trial design of antiangiogenic antibodies. J Natl Cancer Inst 94: 1484 – 1493

Kanehida H, Tamura K, Kurata T, Ueijima H, Nakagawa K, Fukuoka M (2004) Retrospective analysis of the predictive factors associated with the response and survival benefit of gefitinib in patients with advanced non-small-cell lung cancer. Lung Cancer 46: 247 – 254

Korn EL, Arbuck SG, Pluda JM, Simon R, Kaplan RS, Christian MC (2001) Clinical resistance to STI-571 cancer therapy caused by BCR-ABL gene mutation or amplification. Science 299: 876 – 880

Gorre ME, Mohammed M, Ellwood K, Hsu N, Paquette R, Rao PN, Sawyer CL (2001) Clinical resistance to STI-571 cancer therapy caused by BCR-ABL gene mutation or amplification. Science 299: 876 – 880
treatment in patients with non-small-cell lung cancer with postoperative recurrence. *J Clin Oncol* 23: 2513 – 2520

Paez JG, Janne PA, Lee JC, Tracy S, Greulich H, Gabriel S, Herman P, Kaye FJ, Lindeman N, Boggon TJ, Naoki K, Sasaki H, Fujiy Y, Eck MJ, Sellers WR, Johnson BE, Meyerson M (2004) EGFR mutations in lung cancer: correlation with clinical response to gefitinib therapy. *Science* 304: 1497 – 1500

Parulekar WR, Eisenhauer EA (2004) Phase I trial design for solid tumor studies of targeted, non-cytotoxic agents: theory and practice. *J Natl Cancer Inst* 96: 990 – 997

Ross DM, Hughes TP (2004) Cancer treatment with kinase inhibitors: what have we learnt from imatinib? *Br J Cancer* 90: 12 – 19

Rubin BP, Singer S, Tsao C, Duensing A, Lux ML, Ruiz R, Hibbard MK, Chen CJ, Xiao S, Tuveson DA, Demetri GD, Fletcher CD, Fletcher JA (2001) KIT activation is a ubiquitous feature of gastrointestinal stromal tumors. *Cancer Res* 61: 8118 – 8121

Shepherd FA, Pereira J, Ciuleanu TE, Tan EH, Hirsh V, Thongprasert S, Bezjak A, Tu D, Santabarbara P, Seymour L (2004) A randomized placebo-controlled trial of erlotinib in patients with advanced non-small cell lung cancer (NSCLC) following failure of 1st line or 2nd line chemotherapy. A National Cancer Institute of Canada Clinical Trials Group (NCIC CTG) trial. *Proc Am Soc Clin Oncol* 23: 7022

Thomas AL, Morgan B, Drevs J, Unger C, Wiedenmann B, Vanhoefer U, Laurent D, Dugan M, Steward WP (2003) Vascular endothelial growth factor receptor tyrosine kinase inhibitors: PTK787/ZK 222584. *Semin Oncol* 30: 32 – 38

Tokumo M, Toyooka S, Kiura K, Shigematsu H, Tomii K, Aoe M, Ichimura K, Tsuda T, Yano M, Tsukuda K, Tabata M, Ueoka H, Tanimoto M, Date H, Gazdar AF, Shimizu N (2005) The relationship between epidermal growth factor receptor mutations and clinicopathologic features in non-small cell lung cancers. *Clin Cancer Res* 11: 1117 – 1123

van Oosterom AT, Judson I, Verweij J, Stroobants S, Donato DP, Dimitrijevic S, Martens M, Webb A, Sciot R, Van Glabbeke M, Silberman S, Nielsen OS (2001) Safety and efficacy of imatinib (STI571) in metastatic gastrointestinal stromal tumours: a phase I study. *Lancet* 358: 1421 – 1423

Van’t Veer LJ, Dai H, van de Vijver MJ, He YD, Hart AA, Bernards R, Friend SH (2003) Expression profiling predicts outcome in breast cancer. *Breast Cancer Res* 5: 57 – 58

Vogel CL, Cobleigh MA, Tripathy D, Gutheil JC, Harris LN, Fehrenbacher L, Slamon DJ, Murphy M, Novotny WF, Burchmore M, Shah S, Stewart SJ, Press M (2002) Efficacy and safety of trastuzumab as a single agent in first-line treatment of HER2-overexpressing metastatic breast cancer. *J Clin Oncol* 20: 719 – 726

von Bubnoff N, Schneller F, Peschel C, Duyster J (2002) BCR-ABL gene mutations in relation to clinical resistance of Philadelphia-chromosome-positive leukaemia to STI571: a prospective study. *Lancet* 359: 487 – 491

Wedge SR, Ogilvie DJ, Dukes M, Kendrew J, Chester R, Jackson JA, Boffey SJ, Valentine PJ, Curwen JO, Musgrove HL, Graham GA, Hughes GD, Thomas AP, Stokes ES, Curry B, Richmond GH, Wadsworth PF, Bigley AL, Hennequin LF (2002) ZD6474 inhibits vascular endothelial growth factor signaling, angiogenesis, and tumor growth following oral administration. *Cancer Res* 62: 4645 – 4655

Yanagisawa K, Shyr Y, Xu BJ, Massion PP, Larsen PH, White BC, Roberts JR, Edgerton M, Gonzalez A, Nadaf S, Moore JH, Caprioli RM, Carbone DP (2003) Proteomic patterns of tumour subsets in non-small-cell lung cancer. *Lancet* 362: 433 – 439