



Cellular Plasticity and the Untapped Therapeutic Potential in Cancer

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Message from the Guest Editors

Dear Colleagues,

The administration of targeted therapies in cancer patients with well-defined tumour-driving mutations has markedly improved overall survival. Response rates to these therapies, however, remain disappointing, with quantifiable tumour regression limited by the development of acquired resistance. More recent evidence has indicated that targeted therapy can also rapidly induce diverse, genetically-independent, transcriptional programmes resulting in a “drug-tolerant” or “drug persister” cell population. Consequently, during this nongenetic evolutionary phase, cells are able to undergo an adaptive phenotype switch. This cellular or phenotype plasticity exhibited by a subpopulation by tumour cells has been demonstrated to release cells from their dependence on the tumour-driving alteration, resulting in a population of dedifferentiated, slow-cycling cells, capable of surviving continuous drug treatment.





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Message from the Editor-in-Chief

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