Mathematical Modeling of Personalized Androgen Ablation Therapy for Advanced Metastatic Prostate Cancer

Model Aims and Scope
A biochemically motivated model of prostate cancer progression is developed, to conduct a postscriptive analysis of available patient data, for testing the impact of Intermittent versus Continuous Androgen Ablation therapy on the emergence of castrate resistance.

Model predictions are based on fits to Prostate Specific Antigen (PSA) time course data in response to treatment already provided. This results in the identification of new potential biomarkers of disease significance, namely: (i) PSA Velocity; (ii) Cancer cell growth rates; (iii) Cancer cell PSA expression

Key Predictions
• If castration-resistant cancer cells are androgen-independent (positive growth rate irrespective of androgen ablation), both continuous and intermittent therapy will eventually fail.
• Ignoring side-effects, Continuous therapy is preferable to (any) intermittent schedule in general. Intermittent therapy induces earlier onset of castration-resistance.
• Intermittent therapy outperforms continuous scheduling if androgen-independent cells are out competed by androgen-dependent cells.
• If castration-resistant cells are androgen-repressed (negative growth rate in the presence of androgens), intermittent scheduling may delay or prevent the onset of castration-resistance.
• In this case, intermittent scheduling with fixed times on/off androgen ablation outperforms scheduling based on target PSA levels.

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