

**Clinical Trials Summary for out of hours
Important Reference**

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| Acronym study title | Checkmate CA209848 Trial |
| Study Details | A Randomized, Open-Label, Phase 2 Study of Nivolumab in Combination with Ipilimumab or Nivolumab Monotherapy in Participants with Advanced or Metastatic Solid Tumours of High Tumour Mutational Burden (TMB-H) |
| Principal Investigator PI Sub PI's | Dr Dennis Hadjiyiannakis (daytime 01772 522031) Dennis.hadjiyiannakis@lthtr.nhs.uk Dr Pedro Okoh (research medic) (01772 522031) Pedro.okoh@lthtr.nhs.uk Dr Chin (Neuro Oncologist) Chin.lim@lthtr.nhs.uk Dr Appel Wiebke.appel@lthtr.nhs.uk |
| Research Nurse Team | Karen Jones (Karen.jones@lthtr.nhs.uk) Alison Swan (Alison.swan@lthtr.nhs.uk) Nichola Verstraelen (Nichola.verstraelen@lthtr.nhs.uk) Daytime contact (01772 522031) Alternative daytime contact Stephanie Cornthwaite (Stephanie.cornthwaite@lthtr.nhs.uk) (01772 523581) |
| Drug therapy | Randomised to either: Nivolumab monotherapy or Combination of Nivolumab & Ipilimumab |
| In the event that a patient calls this hotline for advise | Please treat as per clinical protocols and inform the clinical and research teams. If you need further specific advice there is a 24 hour line to Bristol Myers Squibb: 01895 523000 where they can give further guidance. |
| Further info | This combination of medications are licensed for use in melanoma currently please refer to these clinical protocols if needed: https://www.healthierlsc.co.uk/application/files/3515/4221/3005/Melanoma_Nivolumab_ipi_v4.pdf |