P-192  HOSPICE DAY THERAPY – A TAILORED APPROACH FOR PATIENTS WITH END STAGE RESPIRATORY DISEASE

1Linda Gregory, 1Julie Davidson, 3Paul Marsden, 1,2Andrew Fletcher. 1St Catherine’s Hospice, Preston, UK; 2Lancashire Teaching Hospitals NHS Trust, UK.

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Background Our hospice currently runs a programme of day therapy for patient with any life-limiting illness. We have previously undertaken a fatigue, anxiety and breathlessness pilot for patients with chronic obstructive pulmonary disease, and work collaboratively with our acute and community trusts supporting a weekly multi-disciplinary team (MDT) meeting to identify patients admitted to hospital potentially in the last year of life, to promote advance care planning discussions. There is an ever increasing emphasis on care which is individualised to patients and their condition(s).

Aims To develop a tailored approach to the support of patients with specific conditions, initially focussing on end stage respiratory diseases, and to demonstrate that this enhances the benefits for patients and their families.

The programme
• 12 month pilot
• 10 week programme of care
• Initial assessment by a member of the hospice palliative care team
• Weekly review by a senior nurse
• Nurse-led follow up clinic, one month after completion with a telephone consultation three months later
• Education and self-management fundamental to the programme. Weekly education sessions for patients and their family/carers, covering management of fatigue, anxiety, breathlessness, nutrition and a session titled ‘understanding your lungs’ to enhance understanding of diagnosis
• Monthly collaborative MDT with acute and community colleagues.

Evaluation The evaluation of this programme will be undertaken throughout and includes:
• Impact on symptom control and quality of life using the Outcome Assessment and Complexity Collaborative toolkit
• Evaluation of education including confidence managing symptoms (patient and family/carers)
• Number of admissions in the 12 months prior to, during and up to six months after commencing the programme
• Number of patients on supportive care registers pre and post programme
• Achieving preferred place of death.

Conclusion Following evaluation of this new approach, we hope to explore if it is applicable to patients suffering other conditions and with robust outcome data, provide evidence for on-going funding.

P-193  ENHANCED DAY HOSPICE – COLLABORATIVE SELF CARE

Donall Henderson, Aine Abbott, Paul McIvor, Damien McMullan. Foyle Hospice, Londonderry, UK.

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Enhanced Day Hospice is a pilot project in Northern Ireland funded by the local Integrated Care Partnership for End of Life Care. This service comprises four designated sessions of medical cover for day hospice including provision of specialist palliative medicine input, working alongside and with support from a staff nurse as well as additional support from the day hospice nursing manager and day hospice volunteers.

It enables a holistic assessment of patients affected by any life-limiting illness. Provided they are well enough to attend, they are also likely to be in the last year (or years) of life, undergoing rapid change in health status, wish to live independently for as long as possible, learning how to manage and minimise the symptoms of their condition(s) with the help (potentially) of family, carers and healthcare staff. Day hospice is a well-established model of outpatient hospice care and was identified as a key aspect of specialist palliative care delivery in the Living Matters Dying Matters strategy for Northern Ireland in 2010.

Patient surveys undertaken across hospices in Northern Ireland indicate that patients greatly value this service and in particular having access to a dedicated day hospice service, whilst carers value the respite time.

Anticipated outcomes of Enhanced Day Hospice:
• Improve ability to manage illness related change for patients, families, carers
• Reduce risks of social isolation
• Anticipate and plan for potential crises
• Improve experience of end-of-life for patients, carers, families
• Reduce reliance on primary and secondary care services during time of intense health challenge.

Other benefits:
• Reduction in hospital deaths
• Improved discharge planning and capacity
• Improved support for Home Care Nursing Team – Increased domiciliary visits
• Improved palliative medicine outpatient capacity
• Reduction in need by patients for other services.

P-194  INNOVATIVE DAY SERVICES OFFER MORE PEOPLE MORE THAN TEA

1Ruth Keeble, 1Katherine Newton, 1Suzanne McArthur, 1Nikki Reed, 1Fiona Dawes, 2Joanne Jordan, 2George Kemohan, 2Kathy Armour. 1Marie Curie Hospice West Midlands, Solihull, UK; 2Ulster University, Newtownabbey, UK.

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Palliative care must change, grow and mature in response to the growing number and complexity of needs of community patients. Such demands particularly challenge conventional delivery of day hospice. In response our hospice redesigned day service provision to modernise bespoke day facilities, appoint a day hospice sister and introduce several novel clinical interventions. Patient outcome measures were introduced and research projects undertaken.

The aim of the study was to evaluate whether extended day hospice services could address more patient needs. Newly developed services were introduced in the period April 2015 – May 2016. Management and clinical key performance indicators were obtained for this period and compared with annual data from the previous year. Two facilitated focus groups were undertaken with the multidisciplinary team and with the management team to
better understand the service developments and participant’s aspirations for day services.

It was found that introducing five additional patient services – Blood transfusion clinic; Progressive Supranuclear Palsy support group; Motor Neurone Disease clinic; Multi System Atrophy support group and a Carers’ support group – led to a 27% increase in attendances. Staff, patients and their families were recruited to three research projects by three hospice staff as a core part of their role, and five staff contributed to qualitative studies or data collection. The qualitative data indicated that staff were pioneering the new well-coordinated services but were challenged by the time available to capture data. The management team had a vision for further expansion of sustainable services and innovative approaches of working in partnership with the local acute trust and community care providers.

In conclusion, a diversification of day services offered to patients and carers can be facilitated with engaged managerial and professional staff without a significant increase in core workforce.

P-195 PALLIATIVE CARE AS AN INTRINSIC PART OF INTEGRATED CARE – DOES THE MODEL FIT?
Caroline Brannan, Nigel Dodds. St Joseph’s Hospice, London, UK
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Introduction Integrated Care(IC) (Ham and Alderwick, 2015) is a model of delivering health and social care, adopted by a number of Clinical Commissioning Groups across East London. This facilitates professionals to work with their external partners, to streamline the patient pathway, and improve patient outcomes. Within this model it is expected that multi-professional teams within the acute and community services, spanning all disciplines, work in collaboration by meeting regularly to proactively plan holistic, person-centred care, rather than focus on separate diseases or co-morbidities, or psycho-social issues, thus aiming to avoid working in silos, in relation to patient care.

Aims One of the main aims of IC is to work towards agreed strategies to avoid hospital admission, and support patients to remain in their own homes. St Joseph’s Hospice community palliative care team are active partners in the integrated care teams, and work with other health and social care providers to ensure patients in the last years of life are enabled to be cared for, and die in their preferred place.

Methods In this presentation we will describe how all relevant professional groups, from all care settings, and involved in patient care meet monthly within every GP practice across the borough in Integrated Care meetings. The focus of these meetings is patient focused, where the most vulnerable patients within the practice are discussed, and a holistic plan of care is agreed using the knowledge and expertise of all the professionals present. This model has replaced palliative care specific meetings where the meeting was focused only on patients approaching the end of life. Through these changes, we will demonstrate the changing nature of referrals to the community palliative care team, which may have implications for the future direction of the hospice.

P-196 CLINIC OR HOME
Linda Gregory, Debbie Bolton. St Catherine’s Hospice, Preston, UK
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Background In September 2012 the increasing number of patients referred to St Catherine’s Hospice Clinical Nurse Specialist (CNS) service, gave the opportunity to explore another option for patients to receive an assessment other than at home. An outpatient clinic service was commenced.

Aims To enable the CNS team to be more responsive to patient referrals. Assessing patients who are well enough in clinic will allow more time for those patients who need to be seen in their own home. A range of other benefits and also some challenges were identified:

- Maintain patient independence
- To optimise patient choice
- It supports the first introduction to a hospice
- Reduce travel costs for CNS team
- Patient may not have transport
- The patient may not feel emotionally able to attend a hospice
- Patients have so many appointments it may be more comfortable for them to be seen at home
- Precedents set by other health professionals that the CNS visits all patients at home
- A change of culture for community nurses who are accustomed to seeing patients at home.

Methods Initially the clinic idea was quite simple in its design; a weekly clinic at the hospice providing four appointment slots.

Results There have been regular audits since 2012 which resulted in further development of the CNS clinics; leading to the current service of two clinics per week. One held at the hospice and a second clinic at the premises of a local cancer charity; providing ten appointment slots per week.

Conclusion It is recognised that patients assessed in clinic are a small percentage of the CNS clinical workload. The aim is now for each CNS to have their own caseload clinic in the community. This is currently being piloted by one CNS having a fortnightly clinic in a GP practice (February 2016).

P-197 THE IMPLEMENTATION OF OUTPATIENT NURSE LED CLINICS BY THE COMMUNITY HOSPICE NURSE SPECIALIST TEAM
Paula Taylor. Wigan and Leigh Hospice, Wigan, UK
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During a Strategic Away Day in November 2013, the Hospice Nurse Specialist Team (HNST) at Wigan and Leigh Hospice identified the necessity to increase responsiveness to patients’ needs during periods of high demand upon the service, by considering ways of increasing overall efficiency. It was agreed to introduce nurse-led clinics for those patients functionally able to attend the hospice, using the WHO performance status tool for guidance.

The HNST have provided face to face assessments in the homes of their patients for many years. Nurse-led clinics was a significant change in approach to the model of care and