

NATIONAL OPTIMAL PATHWAY FOR HEPATOCELLULAR CARCINOMA:

Point of Suspicion to First Definitive Treatment in Adults (aged 16 and over)

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Owner: Upper Gastrointestinal Cancer

Site Group

FOREWORD

The NHS Wales National Optimal Pathways (NOPs) have been developed as part of the Single Cancer Pathway (SCP) programme of work. They aim to establish consistent generic and site specific pathways that describe all routes of entry onto the pathway from the point of suspicion (PoS) of cancer. They describe good practice diagnostic and treatment pathways, the diagnostic pathway, including staging, should be performed within 28 days from PoS; and definitive treatment commenced within 21 days from date of Decision to Treat (DTT). The pathways also describe where patients should receive consistent information and support, tailored to meet their needs.

The NOPs aim to provide a platform to standardise care, reduce unwarranted variation and drive improvement whilst increasing quality across each of the cancer pathways in order to:

- meet the SCP cancer waiting time of 62 days for patients presenting with a suspicion of cancer,
- improve cancer patient experience, and,
- improve cancer patient outcomes throughout Wales to that comparable with the best outcomes in Europe.

The Hepatocellular Carcinoma (HCC) NOP is designed to help HCC cancer service providers and their commissioners see the basic structure of an effective and efficient HCC cancer pathway. In essence, this optimal pathway is about ensuring that each stage of the pathway happens quickly, that communications with patients are effective and that the entire team works in a coordinated but flexible way, focusing always on the patient's journey. It is recognised that the introduction of the National Optimal Pathway for HCC Cancer may present challenges for the Upper Gastrointestinal multidisciplinary teams. However, introducing a nationally agreed, clinically endorsed pathway will support service improvement. They will also provide clarity and consistency for primary care around the referral process into secondary care, including access to diagnostics, to ensure the patients move through the system in a timely manner.

Group Consulted

The Upper Gastrointestinal Cancer Site Group (CSG) is led by Dr. Carys Morgan, Consultant Oncologist and deputy Mr. Nagappan Kumar, Consultant Hepatobiliary Surgeon. The Pathways work has been led by Dr Tom Pembroke, Consultant Hepatologist. The group includes representation from the full range of professions involved in delivering cancer services. They were all able to contribute and comment on the development of the optimal pathway during a range of pathway workshops, CSG meetings and educational days, commenced in September 2019. An early draft was sent to the Upper Gastrointestinal Multidisciplinary Teams (MDTs), Health Boards and Velindre NHS Trust in October 2019. Separate workshops were also held with the Clinical Nurse Specialist (CNS) and Allied Health Professional (AHP) members to embed the Rehabilitation and Person Centred Care (PCC) into the pathway, ensuring all patient's needs are assessed and met in a timely manner. Wider consultation was sought in December 2019 from NHS Wales's stakeholders, including National Imaging and Pathology Networks and the Third sector; 10 responses were received and these were all carefully considered during the revision process, prior to a final version being issued in January 2020.

Dr Tom Pembroke (HCC Lead Clinician for Pathways)

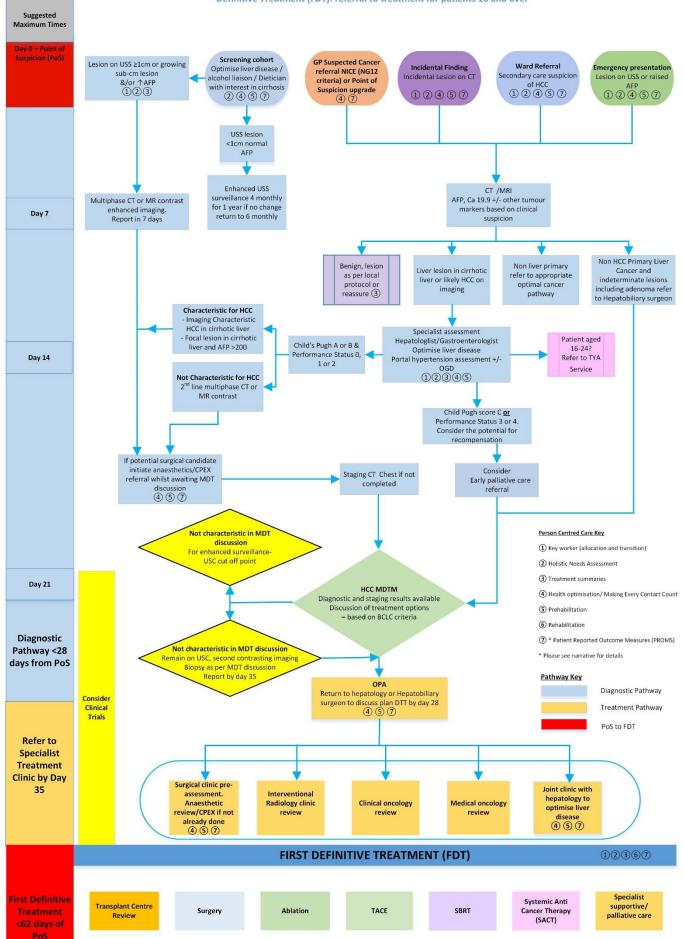
Bethan Hawkes (Macmillan Lead Cancer Nurse - Wales Cancer Network)

Professor Tom Crosby (National Cancer Clinical Director - Wales Cancer Network)

Date 11/02/20



National Optimal Cancer Pathway for suspected and confirmed Hepatocellular Carcinoma (HCC) from Point of Suspicion (POS) to First Definitive Treatment (FDT): referral to treatment for patients 16 and over



DEFINITIONS

Straight to Test	"Following clear referral criteria into secondary care (usually NICE guidance) the secondary care clinician (defined as per local protocol) will arrange a diagnostic procedure as the first episode of care in place of an outpatient episode. The clinician will retain clinical responsibility for the result including acting on the result." Source: Delivering Cancer Waiting Times NHSE
Direct to Test	GPs have direct access to diagnostic endoscopy, ultrasound, MRI, X-ray and CT for people suspected of cancer. Source: National Institute for Health and Care Excellence, Quality Standard 2016 (QS 124)
Safety Netting	"Safety netting is a management strategy of patients, tests and referrals used in the context of diagnostic uncertainty in healthcare. It aims to ensure patients are monitored until signs and symptoms are explained or resolved." Source: Royal College General Practitioners
Decision to Treat	The DATE on which a Decision to Treat is made. For the cancer data sets, the DECISION TO TREAT DATE is the DATE that the consultation between the PATIENT and the clinician took place and a Planned Cancer Treatment was agreed. Source: NHS Data Dictionary
First Definitive Treatment	First Definitive Treatment is the first CLINICAL INTERVENTION intended to manage a PATIENT's disease, condition or injury and avoid further CLINICAL INTERVENTIONS. What constitutes First Definitive Treatment is a matter of clinical judgement in consultation with others, where appropriate, including the PATIENT. Source: NHS Data Dictionary
Teenagers and Young Adults (TYA)	Young people (aged 16-24 years) with cancer have their diagnosis treatment and support agreed and delivered by a cancer-site specific multidisciplinary team and a teenage and young adult multidisciplinary team. Source: National Institute for Health and Care Excellence (NICE), 2014. Cancer services for children and young people (QS55)

CLINICAL EVIDENCE

Point of	Single Suspected Cancer Pathway Definitions (December 2018)
Suspicion/	
Referral	National Institute for Health and Care Excellence (NICE) NG12 Suspected Cancer Referral
	<u>Guidance</u>
	Cancer Risk Assessment Tool (RAT)
	Q Cancer Risk Assessment Tool
Diagnosis and	European Association for the Study of the Liver HCC guidelines 2018
Management	EASL Clinical Practice Guidelines: Management of hepatocellular carcinoma 2018
	https://easl.eu/wp-content/uploads/2018/10/EASL-CPG-Management-of-hepatocellular-
	carcinomapdf
	American Association of the Study of Liver Disease HCC guidelines 2018
	https://www.aasld.org/sites/default/files/2019-06/HCC-Guideline-2018.pdf
	South Wales Hepatocellular carcinomas (HCC) referral and management pathway 2018
	(update due Jan 2020) Not published
	HAP scoring reference
	Kadalayil et al Ann Oncol 2013;24(10):2565-70
	https://www.ncbi.nlm.nih.gov/pubmed/23857958
	<u> </u>
	NICE guideline Radiofrequency ablation of hepatocellular carcinoma 2003
	https://www.nice.org.uk/Guidance/IPG2
	UK Liver transplant selection criteria 2017
	http://odt.nhs.uk/pdf/liver_selection_policy.pdf
	NICE guidance Lenvatinib for untreated advanced hepatocellular carcinoma TA551 2018
	https://www.nice.org.uk/guidance/TA551
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PERSON CENTRED CARE

Key Worker	QS 14: Patients are made aware of who to contact, how to contact them and when to make contact about their ongoing healthcare needs. National Institute for Health & Care Excellence (NICE, 2012) CG138 Clinical Guideline: Patient Experience in adult NHS services
	 A cancer key worker is "a person who, with the patient's consent and agreement, takes a key role in coordinating the patient's care and promoting continuity, ensuring the patient knows who to access for information and advice". 1. All cancer patients must have an allocated key worker 2. Allocation / Review of key worker to take place at key time points including: a. Around the time of diagnosis *please note: allocation of key worker may occur earlier than time of diagnosis, if there is a very high level of suspicion of cancer e.g. evidence obtained via pathology, radiology, endoscopy. b. Commencement of treatment

3. When not under active management by the cancer MDT, the Key worker would be a member of the primary care team. This involvement is critical as many patients living with cancer also live with 2 or more other chronic conditions.

Key workers for cancer patients (Welsh Health Circular /2014/001) GOV.WALES Welsh Government. Cancer Delivery Plan for Wales 2016-2020.

Holistic Needs Assessment and Associated Care Plan

QS 4: Patients have opportunities to discuss their health beliefs, concerns and preferences to inform their individualised care.

QS 10: Patients have their physical and psychological needs regularly assessed and addressed, including nutrition, hydration, pain relief, personal hygiene and anxiety. National Institute for Health & Care Excellence (NICE, 2012) CG138 Clinical Guideline: Patient Experience in adult NHS services

Assessment and discussion of patients' needs for physical, psychological, social, spiritual and financial support should be undertaken at key points including:

- Around diagnosis *please note: undertaking a Holistic Needs Assessment may occur
 earlier than time of diagnosis, if there is a very high level of suspicion of cancer e.g.
 evidence obtained via pathology, radiology, endoscopy; and it is supported by
 professional judgement.
- At commencement, during, and at the end of treatment;

National Institute for Clinical Excellence (NICE, 2004) CSG4 Improving Supportive & Palliative care for adults with cancer

Wales Cancer Network (2016). <u>Guidance document to support implementation of Holistic</u> Needs Assessment (HNA) and associated care plans for the person affected by cancer.

Ideally Holistic Needs Assessments should be undertaken electronically. Welsh Government. Cancer Delivery Plan for Wales 2016-2020. The Macmillan eHNA is *one* tool which is both valid and reliable. Snowden A & Fleming M (2015) Validation of the electronic HNA.

Health Optimisation / MECC

QS 9: Patients experience care that is tailored to their needs and personal preferences, taking into account their circumstances, their ability to access services and their coexisting conditions.

National Institute for Health & Care Excellence (NICE, 2012) CG138 Clinical Guideline: Patient Experience in adult NHS services

Health Optimisation refers to a proactive approach to supporting people who present to NHS services with concurrent comorbid health conditions (e.g. anaemia, diabetes), or health risk behaviours (e.g. smoking, physical inactivity).

Welsh Government (2018) A Healthier Wales.

Welsh Government (2015) Wellbeing of Future Generations Act.

Making Every Contact Count (MECC), is a behaviour change approach that helps health and social care professionals to help people to improve their health and wellbeing through prevention and early intervention. <u>Public Health Wales Strategic Plan 2018-2021</u>

Lifestyle advice / resources are available from Making Every Contact Count (MECC)

*Please note: Whilst addressing concurrent comorbidities and health risk behaviours is the responsibility of **all** health and social care professionals, at **every** contact throughout the pathway; earliest possible intervention **may** impact on cancer treatment choices / outcomes (especially in respect to tobacco smoking). National Institute for Clinical Excellence (NICE, 2018) NG92 NICE Guideline Stop Smoking Interventions and services.

Prehabilitation

Patients should have the opportunity to take part in evidence-based education and rehabilitative activities, including self-management programmes, where available, that promote their ability to manage their own health if appropriate.

National Institute for Health & Care Excellence (NICE, 2012) CG138 Clinical Guideline: Patient Experience in adult NHS services
Welsh Government (2018) A Healthier Wales.

Welsh Government (2015) Wellbeing of Future Generations Act.

Rehabilitation

All patients will have their needs for rehabilitation services assessed, with referral to an appropriate level of rehabilitative support, throughout the patient pathway.

*Please note: not **all** patients will require specialist cancer rehabilitation services. Referral into non-cancer rehabilitation, self-management, and fitness services **may** be suitable to meet some patients' needs.

National Standards for Rehabilitation of Adult Cancer Patients (2010) GOV WALES

National Institute for Health and Care Excellence (NICE) CSG4 (2004) resources for improving supportive and palliative care for adults with cancer.

Patient Reported Outcome Measures (PROMs) / Patient Reported Experience Measures (PREMs)

Patient Reported Outcome Measures (PROMs) are questionnaires that patients are asked to complete **before** and **after** treatment to assess the impact on health and wellbeing.

Some of the Cancer Site Groups (CSGs) have been working with the International Collaboration for Health Outcome Measurement (ICHOM), Value Based Healthcare Team, National PROMS, PREMS, & Effectiveness Programme (NPP&EP) and other partners, to pilot tools and data capture methods, which will inform a consistent approach to PROMS & PREMS for cancer. Welsh Government. Cancer Delivery Plan for Wales 2016-2020.

*Please note: there are outstanding questions relating to tool selection, data capture intervals, data capture methods and data analysis / reporting which **may** prevent the implementation of PROMS & PREMS across all pathways at this time. PROMS & PREMS have been included in the pathways, in recognition of this work, and will be updated pending further advice from the CSGs.

Patient Reported Experience Measures (PREMs) are questionnaires that patients are asked to complete at **any time** during their pathway to help professionals to understand their experience of NHS services. This information is crucial to understanding the value of healthcare as perceived by patients. Welsh Government (2018/19). Chief Medical Officer Annual Report.

*Please Note: Whilst it is good practice to collect PREMS throughout the pathway, there is no current standard for cancer PREMS in Wales; further advice regarding this will be sought via the CSGs in due course.