CervicalCheck Steering Committee

Weekly report to the Minister

24 May 2019

1. Update on support package

The provision of supports to women and families is continuing. In addition, measures have been put in place to ensure that retrospective costs are reimbursed, while an automatic review system is in place to simplify and streamline the claims process to ensure prompt payment of all items covered by the Government decision. As of end-April, €1.42m has been reimbursed in respect of various health and social care costs, €966,000 of which relates to retrospective payments. There will also be additional costs associated with the medical cards that have been issued and the meeting of certain drug costs.

2. Release of records

The HSE remains focused on responding to all requests for records as soon as possible. To aid this process, external legal advisors are liaising with women and their solicitors on the release of slides. The protocol in place ensures the integrity and traceability of slides being transferred; solicitors are required to provide specific information about their chosen laboratory before slides can be released; this ensures the integrity of the slide is protected and all slides can be traced when they leave their current location. The HSE has reported that the average time to deliver slides to independent experts is 28 days. To date, 128 slides out of a total number of 132 have been provided to women and families. A further 4 requests are currently being processed.

3. Independent Expert Panel Review

The protocol for the International Clinical Expert Review led by the Royal College of Obstetricians and Gynaecologists (RCOG) has been published on the Department of Health website.

The HSE project team is holding daily meetings and teleconferences to progress the project. 1,074 (63%) women have consented to take part; the final closing date for consents to be received was Friday 15 February.

The Expert Review Panel has been provided with colposcopy and other data from CervicalCheck in respect of women who have consented to participate, and the transfer of slides from CervicalCheck labs for the purpose of the Expert Panel Review is ongoing. The HSE reports it is continuing to work closely with laboratories to facilitate the transfer. The most recent position is that as of 24 May, approximately 1,566 slides have transferred. Imaging of the remaining slides began on 13 May and is expected to take approximately three and a half weeks to complete. Planning work is ongoing by the HSE in relation to communication of the results of the RCOG review to women. The Information Line remains in service and integrated with the larger helpline, with a low level of calls being received.

Members of the Expert Review Panel reviewed clinical records of women participating in the review in Dublin on Saturday 18th May. The Expert Panel also met with a panel of women and families who have consented to participate in the Review on Friday 17th May. The purpose of this meeting was for the review team to seek a perspective on communication relating to issuing of results, with the aim of enhancing the person-centredness of the reporting process.

4. Smeartaking activity and laboratory capacity

The total number of additional GP consultations was around 112,000. The estimated number of early repeat smear tests which took place between May 1st and December 31st is approximately 57,810, or just over half the number of consultations.

The HSE reports that tests are currently being reported between 3 weeks and 32 weeks of the test being taken. In some cases, this is taking longer. Over half of samples received by the labs are being processed within 8 weeks.

The lab with the largest backlog has ceased accepting new tests from 1 May and will now focus solely on tests in the backlog. The HSE reports that steady progress is being made in dealing with the backlog, with the overall number of outstanding tests reducing. As of 20 May 67,419 smear tests are being processed.

The HSE has agreed with this lab that it carry out a HPV test on smear test samples prior to cytology as a means of prioritising slides appropriately. It is expected that approximately 15% of the total samples taken will be HPV positive. These samples will be prioritised for cytology by the laboratory.

The HSE has advised that it has sourced additional capacity internationally and it is currently working to finalise arrangements in order to incorporate this capacity into the CervicalCheck programme.

5. Introduction of HPV as the primary method of testing

Colposcopy capacity planning is underway by the National Women and Infants Health Programme, which is required to support the introduction of the HPV test. This work includes reviewing current operational pressures for all colposcopy units as well as the impact of the introduction of primary HPV testing and the RCOG review.

A significant volume of work is underway within the HSE to support the introduction of primary HPV screening. A Steering Group is in place to oversee the project, chaired by the Clinical Director of CervicalCheck, with a dedicated project team in place in the National Screening Service to support this work. The negotiations which are underway in relation to the additional laboratory capacity will have an impact on this project and the HSE is working to finalise these arrangements.

6. Colposcopy waiting times

The most recently reported data is valid to end March 2019. 89% of women with high grade abnormalities were seen within 4 weeks of referral (against target of 90%). 90% of women with low grade abnormalities were seen within 8 weeks of referral (against target of 90%).

Currently, time taken in a clinical setting is reported to be considerably longer to facilitate answering questions and putting women at ease, and efforts to manage any impact on waiting times include extra clinical sessions and a focus on waiting list management through appropriate categorisation of referrals.

7. Ex-gratia scheme for non-disclosure

The terms of the CervicalCheck non-disclosure ex-gratia scheme were approved by Government on 11 March 2019, including an Independent Assessment Panel comprising a retired High Court Judge (who will act as Chair), an independent clinician and a person of good standing. The Chair of the Independent Assessment Panel, Mr Justice Aindrias

Ó'Caoimh, was appointed on 5 March 2019, and the remaining two members were appointed by the Minister on 23 April. On 8 May letters issued to women in the 221 cohort, or their next of kin, inviting them to participate in the scheme.

Oversight and engagement with the HSE on modules of its work as follows:

- 1. Management of supports to patients/families
- 2. Provision of documents to patients
- 3. Interface with RCOG Review
- 4. Management of laboratory capacity issues
- 5. Introduction of HPV Screening
- 6. Colposcopy

1. Management of supports to patients/families

Significant Issues

There are no exceptional items to report in relation to Community Supports.

2. Provision of documents to patients

Significant Issues

Current position, issues & challenges

The team remains focused on responding to all slide requests as soon as possible - the average time to deliver slides to the independent expert is 28 days. Weekly operational meetings continue to monitor the laboratories.

The HSE has provided 128 slides out of a total number of 132. There are 5 currently being processed.

A total number of 574 records have been provided, out of a total number of 584 requests. There are 10 outstanding which are being processed.

Issues: The HSE has identified a significant number of cases where it has not been informed of the requestors name or/and their designated lab expert where the slides are to be sent to – as a result we are working with those solicitors to ensure that all required details are being sent to the labs.

Project Governance	 Support Team continues to hold daily meetings and teleconferences to progress deliverables identify critical actions / timeframes, areas for escalation, and project RAIDS (Risks, Actions, Issues, Decisions). Structures and processes are being established to support disclosure of results with reference to existing processes already documented e.g. HSE Open Disclosure Policy, Safety Incident Management Policy, Lookback Review Guidance, etc. 								
Patient Support Services		Action	Activities Planned						
	• 1,074 (63%) Panel Revie	women have cow.							
Laboratory Logistics	Lab	Number of slides requested to date 340	Number of slides sent to RCOG- relating to slide requests 14	Total number of available slides remaining to be sent to NBHT 310	 Maintain regular contact to Medlab and the Coombe on imaging of Medlab slides, to 				
	QUEST Coombe	274 1,088 62	265↑4 1,027↑11 62↑ 1	2 5 0	ensure slides are imaged and transferred as per proposal agreed.				
	Total	1,764	1,368 ↑1 6 (78%)	317 (18%)					
	 16 slides transferred from 3 labs in the last week. The Coombe has transferred all requested slides to the RCOG, with 0 outstanding. Proposal received from Medlab on Tuesday 8th May on the imaging of requested slides at the Coombe, to commence May 13th. It is expected to take at least 3 ½ weeks to image all requested slides. Proposal agreed by the Coombe, and shared with RCOG and DOH. 								
Information Services	There were	<u> </u>							
Case Management System (CMS)	oversee the members of	A Project Manager has been assigned from Office of the CIO to oversee the maintenance and development of the CMS – met members of the HSE RCOG Support Team on 8 th May to discuss planning and requirements for CMS.							
Acute & Community Services	The Expert Review Panel identified a list of women for whom clinical records are required for review when they are in Dublin on Saturday 18 th May. In line with the agreed SOP with acute services, this list has been separated by hospital and circulated to the appropriate hospital group for retrieving the relevant files. Files are due for receipt W/C May 13 th .								
	the Review Expert Revie request the issuance of	vomen (or their has been forme ew team on Frid group's views or results, with the sof the reportin							
Clinical Open Disclosure	of results w	and processes ar ith reference to en Disclosure Po							
	sent letter o	commissioner of invitation and don Wednesda	0,						
Current position, significant issues	The CervicalCheck contracted laboratories have identified a number of slides that are currently unavailable for the RCOG Review, as they were previously released to an independent expert reviewer and have not yet been returned to the originating laboratory as yet. Letters are issuing from the NSS to these women and their solicitors where relevant. Unfortunately, if those slides are not received by the expert panel by 30th April, they will be unable to review those cases.								
				eve also identified a etrieve these slides.	number of slides that are proving difficult to				

^{*} The total number of slides sent to the UK may be greater than total number of slides requested from labs due to troubleshooting process whereby 2 slides (original and treated) are prepared from one sample

4. Management of Laboratory Capacity Issues						
Uptake of Smear	Out of Cycle Smears					
Tests	The total number of additional GP consultations between May 1 st to December 31 st was more than 112,000. The estimated number of early repeat smear tests to take place in the period of May 1 st to December 31 st is in the region of 57,810.					
Average Time for Processing Results	We remain extremely concerned at the length of time being taken for reporting of cervical smear tests, which regrettably are being reported between 3 weeks and 32 weeks of the test being taken. In some cases this is taking longer. However, it is worth noting that over half of samples received by the labs are being processed within 8 weeks.					
	We have worked with existing private providers, other private providers and public service providers in other countries to try and grow our laboratory capacity. Some of our existing providers have managed to reduce the wait times and we continue to work with others to try and find additional capacity. While we continue to pursue active leads this has proved very challenging due to the global shortage in cytology. This has been caused as a result of the reduced cytology requirement as countries implement HPV primary screening – which sees a reduction of c80% for cytology requirements. We are actively trying to identify possible solutions that will help reduce the wait times which we know are causing a lot of anxiety for women.					
	Whilst this is very undesirable, our clinical advice is that this poses a very low risk to women. Notwithstanding this, we recognise that these delays are extremely difficult for women and we are making every effort to improve this situation. We have made significant improvements in the turnaround times with two of our three laboratories and are working closely on an improvement plan with the third laboratory. We are absolutely focused on reducing waiting times for results as quickly as possible.					
	The HSE has been working closely with the lab with the largest number of tests waiting to be processed, MedLab Pathologies Ltd, to clear all outstanding smear tests as a matter of priority. This lab processes smear tests for women based mainly in the south and west of the country. We are making some progress in this regard; with the overall number of outstanding smear tests steadily reducing.					
	This work with MedLab Pathologies Ltd is being done based on a HPV initial testing model; i.e. HPV testing will be carried out on smear test samples prior to cytology.					
	While the human papillomavirus (HPV) is a very common virus and usually clears without treatment, some types can cause changes in the cells of the cervix that can later develop into cervical cancer. This HPV testing model was chosen as the most effective way for the lab to process all outstanding tests based on prioritising women most at risk.					
	This means that: - All outstanding tests will have an initial HPV test carried out Tests which report as HPV positive will be prioritised for cytology, as these women are considered to have the highest clinical needs Tests which report as HPV negative will have cytology performed as a second priority, as these					
	women are considered to have lower clinical needs. While this model may result in an initial further delay for women with negative HPV results, it will ultimately allow women with HPV positive results, who are considered most at risk, to be prioritised					
	and for those women to receive necessary follow-up care. We hope that this process will provide reassurance to women whose outstanding tests are with MedLab Pathologies Ltd that we are doing out utmost to expedite these tests.					

HPV Steering Group established with NSS, HSE and service user representatives Project team established with Continue Charles NSS and USE membership.																
Project team established with CervicalCheck, NSS and HSE membership Detailed project plan on Project Vision managed by PMO team.																
Detailed project plan on Project Vision managed by PMO team Project to any act blish admirblish admirblish density to any act and work stream leads.																
Project team established with identified project manager and work stream leads Continued the Clinical Directors assume and 4 (22).																
 CervicalCheck Clinical Director commenced 4/02 National laboratory QA lead appointed. Commenced 14/1 Colorscopy load still outstanding. There is colorscopy representation on the Clinical Advisory 																
										 Colposcopy lead still outstanding. There is colposcopy representation on the Clinical Advisor Group (CAG). Meetings with colposcopists are held regularly. 						
Stabilisation of current programme and capacity planning- increase in laboratory test																
volumes in 2018 has resulted in significant lengthening of the process and reporting timeline																
Capacity planning is underway to address the backlog with a detailed planned impact assessment and options appraisal. • Public confidence- reporting times and retests are impacting on confidence in the cervical screening service.																
									Procurement- despite on-going work to develop services in the Coombe to maximise public							
									provision in the future, this work is time dependent. Additional lab services will be required							
									for the HPV primary screening transition as there is not sufficient capacity available in the							
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6. Colposcopy

- CervicalCheck has established a network of quality assured colposcopy clinics for women requiring further investigation following a smear test. A woman can be referred to one of 15 colposcopy clinics located nationwide.
- Extra clinical sessions have been added to reduce waiting lists
- Within the current climate time taken in a clinical setting is considerably longer to facilitate answering queries and putting women at ease
- Extra efforts made when appointments are cancelled to fill the vacant slot to further reduce waiting lists.
- Extra efforts to ensure the increased referrals are categorised in a prompt manner to ensure high and low grade are seen within guidelines

Colposcopy data

March					March Data		
	Monthly		Annual YTD		Month Year Colposcopy Clinic (& associated histology laboratory)	Average (Combined)	
	Projected	Actual	Projected	Actual	*Waiting time HG end month - Target 90% to be seen within 4 weeks of referral	89%	
Referrals	1,625	1,515	4,875	4,486	*Waiting time LG end month - Target 90% to be seen within 8 weeks of referral	90%	
					*HG - High Grade, LG - Low Grade **Figures for the Coombe not available this month		