

## Discussion outputs from DIGITAL HEALTH LEADERSHIP SUMMIT, March 2021

### Topic 18: How do we deliver a better patient experience through digital?

This topic was discussed by a group in Auckland.

#### Auckland delegates' responses

<b>Definition</b>	<ul style="list-style-type: none"><li>• Safe and comfortable</li><li>• Digital triage for items and issues for doc prep</li><li>• Clinical and care for</li><li>• Listened to</li><li>• Questions answered</li></ul>
<b>Goals</b>	<ul style="list-style-type: none"><li>• Preserving the great things about seeing your doctor</li><li>• Keeping it personal and caring</li><li>• Feeling of personal care and attention</li><li>• Extended patient carers (family etc.)</li><li>• Closing the loop in a timely manner</li><li>• Patients being empowered to drive</li><li>• Cost-effective service</li><li>• Patient experience more inclusive of whanau</li><li>• Patient ownership</li><li>• Ask what experience they want</li><li>• What outcome do you want?</li><li>• Health providers to collaborate for me. GP, specialist, physio (tack!)</li><li>• D.R.Y. principle</li><li>• Choice!</li><li>• The experience begins early when they start looking</li><li>• Knowing the right process of seeing clinician?</li><li>• Less waiting time</li></ul>

	<ul style="list-style-type: none"> <li>• I want to live longer, with a better quality of life</li> <li>• Own my own data. It should travel with me</li> <li>• I understand the information and can update it</li> <li>• Clarity of process. Opportunity to give feedback. Listen to me!</li> <li>• I want to update my information</li> <li>• Manage my own appointments</li> <li>• I've got the best working for me</li> <li>• Continuous record of care – birth to death (continuity of care)</li> <li>• Patient owns their information and shares it with caregiving parties</li> <li>• Single source</li> <li>• Measure patient experience over time – CX</li> <li>• See “patients” as “consumers” and measure metrics</li> <li>• Reduce duplication</li> <li>• Reduced time in front of clinician – self-managing</li> <li>• Information follows me in my healthcare journey</li> <li>• Integrate all social services in and outside of health system</li> <li>• Have patients involved more up-front (engagement)</li> <li>• Embrace indigenous models of healthcare (holistic)</li> <li>• Equitable outcomes and experiences for everyone (success is individual)</li> <li>• Life coaching model. Capabilities, strengths, gap, how to help gap → track progress as you go – feedback loop → mentoring for each individual from childhood (feedback)</li> </ul>
<b>Barriers/challenges</b>	<ul style="list-style-type: none"> <li>• Appointment: <ul style="list-style-type: none"> <li>○ ✓ appointment</li> <li>○ ✓ prescription</li> <li>○ ? the actual appointment</li> </ul> </li> <li>• Older demographic – tech?</li> <li>• Incentives?</li> <li>• No incentives to improve</li> <li>• Measurement</li> </ul>

- Staffing issues – digital to help?
- Aging GP population?
- Waiting room! ... 15 minutes ...
- Increase “the 1 hour’ per year
- Use technology to recreate and elevate the experience
- Pushing admin jobs to them (clinicians)
- Right access
- Patients must own their own knowledge and experience
- Controlling the right information, dissemination
- There’s no urgent need to drive P.X.
- Changes: barriers to access/learning
- Better connectivity
- Upskilling clinicians on digital
- Lack of access to records
- Cultural - healthcare system
- Clarity of clinician – start → stop?
- Access to preventative, information, etc.
- Siloed nature of healthcare industry
- Funding under-resourced
- Funding doesn’t follow the patient
- Census completion!
- Funding structure is out of date
- Not measured – lack of KPIs
- Health literary and access to technology
- Getting care – I want where I want, when I want it
- Getting care out of hours – my hours
- Siloed nature of care – repetition – no code sign-on at systems level
- Ask “what matters to me” (not just what’s wrong with me – outputs)
- Output rather than outcomes focused

	<ul style="list-style-type: none"> <li>• 'To' rather than 'for' 'with'</li> <li>• 'outputs' (metrics) are not 'outcomes'</li> <li>• Health system is not built from a consumer's experience. Still on post WWII model</li> <li>• Current system rewards duplication</li> <li>• Patient 'experience' shouldn't be what we're gunning for. Should be 'patient engaging them'</li> <li>• ACC funding is based on <u>more</u> cases = more funding (no incentive to solve problems)</li> <li>• Current talk about 'mind/body and 'wellness' is new</li> <li>• System focused on 'disease' rather than 'wellness'</li> <li>• A portion of population limited access to digital services (geographical, age, socio-economic)</li> <li>• Many people do not want to take responsibility for own healthcare</li> </ul>
<b>Solutions/ideas</b>	<ul style="list-style-type: none"> <li>• Questions answered, empathy</li> <li>• Regular low-fi check-ins</li> <li>• Simple, remote monitoring</li> <li>• Create feedback loops. Measure the experience</li> <li>• Families working together better</li> <li>• Patient journey + notifications</li> <li>• Free 'open source' technology</li> <li>• Automated notifications, feedback</li> <li>• Opportunities for improving digital literacy</li> <li>• NPS standardised</li> <li>• Doctor "traffic" and triage app</li> <li>• Matching demand with supply</li> <li>• Set your own goals/outcomes app</li> <li>• Health wallet \$\$</li> <li>• Cross regions</li> <li>• Giving feedback!</li> <li>• Shared access to medical records – across departments, clinics. Support person or family member access as well.</li> <li>• Health prevention – as a service</li> <li>• Whole view of my healthcare: Provider and end user</li> </ul>

- Giving patients a voice
- Digital literacy – whose responsibility is to it meet a certain ‘level’ of literacy
- Patient needs to repeat themselves
- Siloed information, e.g., GPs, hospitals
- Platforms and systems not talking to each other
- Getting notes, history
- Should patients have the right to add to their notes? → ManageMyHealth is doing some of this
- Often, we don’t get correct or full patient history. Need one place, on history
- Cohorts not able to release information
- Legal restrains for sharing information
- Privacy laws, legislation
- Old technology
- Test and other data, e.g., prescriptions, drug issues, not shared between regions
- Not all DHBs have systems – still quite paper-based
- A lot of information is in PDF format and can’t extract/search the information
- Tertiary institution (e.g., university) care is unpaid
- GP to GP transfer data goes into a central database, but not from DHB
- Funding does not follow the patient (e.g., from hospital to GP)
- Pushing care out to the community is not funded
- Admin overhead
- What do patients think?
- Patients think that systems talk to each other, but they don’t
- Ministerial responsibility – where do the \$\$ come from?
- Central repository of data (single source of truth)
- Lifetime history of medical care
- Self-service style of digital channel
- Patient able to determine when they receive care
- Systems integration
- Portal? App? Cloud-based

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|  | <ul style="list-style-type: none"><li>• Fit for purpose system</li><li>• Reduce # - DHBs to reduce duplication</li><li>• Apps and teleconsultation are leading the way</li><li>• To be empowered, need access to information</li><li>• If single source not feasible, then let the consumer be the system – integrated</li><li>• Invest \$ in helping people access digital services</li><li>• A design that is seamless and feels like “magic”</li><li>• Already a wealth of wellness apps – could bring all information together</li><li>• Encourage people to live their best life (would people even know what that would be?) – people who would be most likely to do this area are already doing well in health system</li><li>• Improving the health system may not be an ‘exiting’ digital solution. Low tech engagement</li><li>• Co-design solution with communities</li><li>• Encourage ‘hope’</li><li>• Digitally enabled health advocacy/literacy</li><li>• Digital tool to help community connection, advocacy</li></ul> |
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