

The Hong Kong Principles for Assessing Researchers: Fostering Research Integrity

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29 **Abstract**

30

31 The primary goal of research is to advance knowledge. For that knowledge to benefit research and
32 society, it must be trustworthy. Trustworthy research is robust, rigorous and transparent at all stages of
33 design, execution and reporting. Initiatives such as the San Francisco Declaration on Research
34 Assessment (DORA) and the Leiden Manifesto have led the way bringing much needed global attention
35 to the importance of taking a considered, transparent and broad approach to assessing research quality.
36 Since publication in 2012 the DORA principles have been signed up to by over 1500 organizations and
37 nearly 15,000 individuals. Despite this significant progress, assessment of researchers still rarely includes
38 considerations related to trustworthiness, rigor and transparency. We have developed the Hong Kong
39 Principles (HKPs) as part of the 6th World Conference on Research Integrity with a specific focus on the
40 need to drive research improvement through ensuring that researchers are explicitly recognized and
41 rewarded (i.e., their careers are advanced) for behavior that leads to trustworthy research. The HKPs
42 have been developed with the idea that their implementation could assist in how researchers are
43 assessed for career advancement with a view to strengthen research integrity. We present five
44 principles: responsible research practices; transparent reporting; open science (open research); valuing
45 a diversity of types of research; and recognizing all contributions to research and scholarly activity. For
46 each principle we provide a rationale for its inclusion and provide examples where these principles are
47 already being adopted.

48 **Introduction**

49 In a quest to advance knowledge, researchers publish approximately 1.5 million journal articles each
50 year. The presumption is that this literature can be used by other researchers, stakeholders, and the
51 wider society because it is trusted, robust, rigorous and complete.

52
53 The approach taken to validating research and its outcomes differs depending on the nature of the
54 research. For example, to rigorously examine the effects of a health intervention, trial participants
55 (human or animal) are typically required to be randomized to the intervention being studied. Many
56 researchers advocate registration of protocols as a way to ensure transparency and to enable others to
57 engage with their research. Subsequently, the use of reporting guidelines can help ensure complete and
58 transparent reporting of the researchers' methods and results. When the research is being
59 disseminated, the research team would ensure that the associated data, materials and any analytical
60 code are made available as an integral part of publication. Such data sharing facilitates re-analysis of the
61 data to check reproducibility and to perform secondary analyses.

62
63 Although some mechanisms exist to support researchers in ensuring transparency at all stages of design,
64 execution and reporting, there is not widespread adoption of these practices. There are many
65 interwoven reasons for this. One contributing factor, we argue, is that little emphasis is placed on the
66 rigor of research when hiring, reviewing and promoting researchers. Working together across the
67 research sector as a whole to address this systemic issue, we believe, offers a global opportunity to
68 improve research and impact.

69
70 We developed the Hong Kong Principles (HKPs) as part of the 6th World Conference on Research
71 Integrity (WCRI) specifically to drive greater recognition for researchers who commit to robust, rigorous
72 and transparent practices (i.e., their careers are advanced) (see Figure). If implemented, the HKPs could
73 play a critical role in evidence-based assessments of researchers and put research rigor at the heart of
74 assessment, as well as open up research to the wider benefit of society.

75
76 We propose five principles, each with a rationale for its inclusion. We illustrate these principles with
77 examples where we know they exist. These examples are not exhaustive, and many are relevant to more
78 than one principle. Together, they illustrate of a breadth of approaches as to how these principles can
79 operate at the very highest levels of international research.

80

81 Early drafts of the HKPs were circulated to the 700 participants registered for the 6th WCRI. Further
82 discussions took place during two sessions at the 6th WCRI. A penultimate version was uploaded on the
83 6th WCRI website after the conference. More than 100 people provided input and feedback. We
84 acknowledge all of these valuable contributions and the global leadership of those working on the San
85 Francisco Declaration on Research Assessment (DORA), the Leiden Manifesto and other initiatives to
86 promote the responsible use of metrics, which have laid the foundations for much of our work (1-4).

87

88 **Principles**

89 **Principle 1: Assess researchers on responsible practices from conception to delivery, including the**
90 **development of the research idea, research design, methodology, execution and effective**
91 **dissemination.**

92

93 *Rationale*

94 The numbers of publications, citations, and total volume of grants are often still the dominant metrics
95 used by research institutions for assessing and rewarding their researchers (1-4). Providing bonuses to
96 academics for publishing in certain journals (i.e., merit pay) is also common in many parts of the world
97 (5-7). These assessment criteria tell assessors little about the researchers and the rigor of their work;
98 thus they are not particularly 'responsible' metrics. These metrics can also be unduly influenced by field
99 and citation practices and provide little information about a publication's (and therefore a researcher's)
100 contributions to research and society. Other criteria are required to provide a broader view of markers
101 of best practices: for example, the extent to which a researcher develops research questions with the
102 involvement of appropriate members of the public (see Figure).

103

104 *Current implementation*

105 The Canadian Institutes of Health Research's Strategy for Patient-Oriented Research (SPOR) is a multi-
106 million-dollar initiative to bring patients into a broad range of activities regarding research across
107 Canadian provinces and territories (8). Patients are now active in the development of research projects
108 in setting priorities and formulating study questions. The Ontario response (Ontario SUPPORT Unit) has
109 included a series of articles with patients taking a leadership role in co-authoring the content (9). In the
110 UK, the James Lind Alliance, funded by the UK National Institute of Health Research (NIHR), is a

111 successful example of including patients, carers and clinicians to develop priority-setting partnerships
112 (10) and question formulation (11).

113

114 With a focus on enhancing reproducibility the US National Institutes of Health (NIH) have revised their
115 application instructions and review criteria to strengthen scientific rigor and transparency (12). One of
116 the resources they recommend is the Experimental Design Assistant (EDA) developed by The National
117 Centre for the Replacement, Refinement & Reduction of Animals in Research (NC3Rs). This 10-module
118 online tool was developed to assist researchers in the design and analysis of animal experiments. It
119 includes dedicated support for randomization, blinding and sample size calculation. It can also be used
120 to help researchers prepare the experimental design information and analysis plan requested for grant
121 applications (13). The NC3Rs encourages the use of the EDA so that applicants can generate an EDA
122 report, which can be submitted in place of the experimental design and methodology appendix.

123

124 Other examples of preferred criteria include social media metrics as indicators of disseminating research
125 (14), public lectures about the results of a research project, public engagement and other types of
126 events that bring together funders, researchers and other stakeholders to work on an effective
127 communication plan of the research program (15). Organizations such as the Wellcome Trust are taking
128 a holistic attitude to redefining their approach to engagement explicitly to help people feel empowered
129 to access, use, respond to and create health research (16).

130

131 **Principle 2: Value the accurate and transparent reporting of all research, regardless of the results.**

132

133 *Rationale*

134 Failure to publish all findings of all studies seriously distorts the evidence base for decision making. For
135 example, a systematic review of trials of reboxetine for treating depression found that almost three
136 quarters of included patients were in unpublished trials (17). Selective publishing of research with
137 positive results (i.e., publication bias) distorts science's evidence-base and has been demonstrated in a
138 variety of disciplines including economics, psychology and clinical and preclinical health research (e.g.,
139 18). Furthermore, the frequency of other reporting biases (e.g., switched primary outcomes without
140 disclosure, and spin) is around 30% (19). This is unacceptably high and diminishes the trustworthiness
141 and integrity of research (7). It also appears that Promotion and Tenure Committees (PTCs) generally do

142 not give sufficient importance to registering protocols and data analysis plans, full publishing of
143 completed studies or making data, code, and materials available (20).

144

145 *Current implementation*

146 Study registration and reporting guidelines are useful tools to help improve the completeness and
147 transparency of a very broad spectrum of research (e.g., 21-24). As part of the editorial policies of the
148 Wellcome Trust’s open access publishing platform (Wellcome Open Research (WOR)), authors are
149 required to use reporting guidelines for protocols (e.g., SPIRIT) and completed studies (e.g., ARRIVE
150 (25). Other funders, such as Gates Open Research (26), the NC3Rs Gateway (27) and the Association of
151 Medical Research Charities (28), do likewise. To help reduce publication bias, WOR also requires
152 registration through one of several different options (e.g., registered reports) (25). Similarly, to promote
153 the registration and publication of all research the NIHR in the UK indicate that “When submitting an
154 application to NIHR programmes for funding for a new clinical trial, the applicant must disclose past
155 publication and trial Registration history for any relevant publications and research grants held,
156 referenced in the application.” (29). While these are examples of best practice from funders, we were
157 unable to find any research institution that has incorporated them into researcher assessments (20).

158

159 Several research institutions (e.g., University of Toronto) are now recommending that their clinical
160 trialists use SEPTRE (30), a web-based protocol creation and management tool. When SEPTRE is used,
161 protocol information for trials is automatically registered in clinicaltrials.gov. This saves time and helps
162 the researchers, and their research institutions, to maintain best publication practices (e.g., trial
163 registration). Some journals in the social sciences, particularly psychology, use registered reports to help
164 ensure that research is published regardless of its results (31,32).

165

166 **Principle 3: Value the practices of open science (open research) - such as open methods, materials and** 167 **data.**

168

169 Rationale

170 Openness in research is more than just access to research – it brings equality to the research process. It
171 encompasses a range of practices across the entire lifecycle of research (33). Access to research should
172 not be about who has the resources to pay to see behind a paywall, typically subscription journals.
173 Healthcare and social policy decisions should be made based on access to all research knowledge rather

174 than only a part of it (34). A considerable amount of public funds is used for research and its results can
175 have profound social impact. Preclinical scientists are committing to openly share their laboratory
176 notebooks (35) to streamline research, foster collaborations and reduce unnecessary duplication. In an
177 effort to deter questionable authorship practices, the Consortia Advancing Standards in Research
178 Administration Information supports the CRediT taxonomy (36) as a way for research authors to more
179 openly describe how each person has contributed to a research project.

180

181 Data sharing is another example of openness but is not common practice in clinical research (with some
182 exceptions, such as genetics) (37) although patients seem supportive of sharing their data, at least of
183 randomized trials they have participated in (38). Data sharing is also not considered standard in many
184 other disciplines. Without data sharing it is difficult to check the selectivity of reports; data sharing is key
185 to addressing the reproducibility crisis (39) and building trust (40). There are varying estimates as to
186 which proportion of research is made available through open access mediums, such as open access
187 journals, repositories, or as preprints, but it is far from 100% (41).

188

189 *Current implementation*

190 Ghent University, Belgium, has employed data sharing guidance stating, "Sound data management is a
191 basic requirement for this (academic analysis) and provides additional guarantees for a flawless
192 methodology, for sharing, and reusing data by other researchers in an Open Science context and for the
193 accountability of a researchers own academic integrity" (42). The Nanyang Technological University
194 (NTU), Singapore, implemented an Open Access policy in 2011. All NTU faculty and staff must deposit
195 their final peer-reviewed manuscript of journal articles and conference papers in the Digital Repository
196 (DR-NTU) maintained by the Library upon acceptance of their publications. At NTU's faculty of medicine,
197 random data audits are conducted on the submitted (required) Data Management Plans (DMPs) and
198 checks are made to see if the final data are indeed shared on NTU's open access data repository DR-
199 NTU.

200

201 To help facilitate data sharing the University of Cambridge has introduced the concept of 'data
202 champions' (43). Here, volunteers advise members of the research community on proper handling of
203 research data supporting the use of the Findable, Accessible, Interoperable, and Re-usable (FAIR)
204 research principles (44). Delft University of Technology, The Netherlands, has taken this concept a step
205 further and implemented it as a career assessment criterion (45). The University of Glasgow's academic

206 promotion criteria explicitly allows for data sharing as a research and scholarship output (to support
207 replication) (46).

208
209 Some journals have also established strong data sharing policies. For example, the PLOS journals
210 “require authors to make all data underlying the findings described in their manuscript fully available
211 without restriction at the time of publication. When specific legal or ethical requirements prohibit public
212 sharing of a dataset, authors must indicate how researchers may obtain access to the data. Refusal to
213 share data and related metadata and methods in accordance with this policy will be grounds for
214 rejection.” (47). Given that societal benefit is part of an emerging career assessment, clinical researchers
215 should also respond to a growing view that patients want their data shared (38).

216
217 Open research is supported by key infrastructure compliance, such as requiring an Open Researcher and
218 Contributor ID (ORCID) by every researcher, whereby each researcher can be uniquely identified. A
219 recent letter from global funders committing to Implementing ORCIDs for all researchers is a significant
220 step forward (48). This was recently implemented at the Ottawa Hospital Research Institute (49). In
221 Australia and New Zealand there is a consortium that supports ORCID nationally.

222
223 The NIH promotes the use of preprints in grant applications (50) as do all major UK public funders (e.g.,
224 Medical Research Council, UK) (51), The Wellcome Trust made them compulsory for work in health
225 emergencies and promotes their use widely in particular for early career researchers (52).

226
227 **Principle 4: Value a broad range of research and scholarship, such as replication, innovation,**
228 **translation, synthesis, and meta-research.**

229
230 *Rationale*

231 A system that rewards benefit to society and encourages trustworthy and important research needs to
232 take the different types of research into account: creating new ideas; testing them; replicating key
233 findings; synthesis of existing research; developing and validating new tools; measures or methods; etc.
234 Different indicators and criteria need to be developed that are relevant to these different types and
235 stages of research (see Figure). This includes different timeframes of assessment for different types of
236 research.

237

238 Incentives that encourage one fixed idea of the ‘right kind’ of research will be slow, or even stall,
239 progress. So-called blue-sky research that builds on chance findings or curiosity-driven research based
240 on ‘out-of-the-box’ thinking should be possible and encouraged, as well in an academic reward system
241 that values societal progress (53). For example, the discovery of graphene at the University of
242 Manchester, UK, was the result of Friday afternoon discussions outside the ‘normal’ research activities
243 (54). Other examples from a broad range of disciplines exist (55). The short-term nature of academic
244 reward cycles makes this kind of research less attractive for funders, institutions and individual
245 researchers. Equally, replication studies or research synthesis efforts are often not regarded as
246 innovative enough in researcher assessments despite their critical importance for the credibility of
247 research, or for a balanced and robust systematic presentation of all available evidence, respectively
248 (39,56). This is not universally appreciated by PTCs. Research on research and meta-research are
249 practiced at, for example, at METRICS (Stanford, USA) (57), QUEST (Berlin, Germany) (58) whose focus is
250 on clinical and preclinical meta-research, and the Meta Research Center at Tilburg University (59)
251 (Tilburg, The Netherlands) whose focus is on the social sciences. Such activities are important to inform
252 and improve research practices and therefore contribute to making research more reliable and relevant.

253

254 *Current implementation*

255 Some funders have already recognized the relevance of a broad range of research activities. The
256 Research Impact Assessment Platform (Researchfish) works to capture some of this diversity and can
257 generate reports on the impact of a broad spectrum of funded research (60). The Wellcome Success
258 Framework highlights the importance of a long-term vision and shared objectives in order to take a
259 more balanced approach to assessment (61). The German Federal Ministry of Science and Education is
260 funding preclinical confirmatory trials (62).

261

262 The Wellcome Trust has developed a new Longitudinal Population Studies Strategy, funded data re-use
263 prizes (63) and supports research on research (64). All approaches are aimed at valuing a broad range of
264 scholarship and maximizing the value of research. The Netherlands Organization for Scientific Research
265 is in its third call for replication studies (65). Research on research and meta-research are also gaining
266 momentum and now have some formal outlets. For example, PLOS Biology and eLIFE have a meta-
267 research section in their journals (66,67). We were unable to find any academic institution that has
268 incorporated replication or meta-research into their career assessment portfolio (20). NIHR requires the
269 completion of a systematic review prior to funding any new research (68). The NC3Rs have also

270 promoted the importance of systematic reviews for providing a rationale for project proposals (69,70).
271 In the event that such a review does not exist, they provide funding to perform one.

272

273 **Principle 5: Value a range of other contributions to responsible research and scholarly activity, such as**
274 **peer review for grants and publications, mentoring, outreach, and knowledge exchange.**

275

276 *Rationale*

277 As discussed alongside Principle 1, research assessments frequently focus on a narrow range of easy to
278 measure metrics including publications, citations and funding income (1,20). For the research ecosystem
279 to function optimally, other research activities are also essential. Peer review remains the cornerstone
280 of quality assessment of grants, publications and conferences. The quality of peer review contributions
281 to journals and funders, should also be part of assessments for promotion and tenure as should
282 contributions to various research infrastructure, oversight, or regulations. Equally, contributions to
283 improvements that go beyond an individual-centered approach for assessment should be considered.
284 These activities are currently largely missing from PTCs (20). Contributions to developing the careers of
285 others at all stages of their career is critical as are contributions various committees related to research
286 (e.g., assuming the role of an editor). How best to do this without creating further barriers and
287 bureaucracy, however, has long been debated (71).

288

289 Any reward system that has the whole research enterprise at heart and aims to foster a climate
290 conducive to trustworthy and useful research with the highest regard to integrity, needs to find ways to
291 incorporate these vital roles into its overall assessment structure.

292

293 *Current implementation*

294 Macquarie University, Sydney, Australia, has some exciting initiatives in their new academic promotion
295 policy which includes five pillars one of which is in leadership and citizenship. Here researchers can show
296 their alignment with the university's values and broader contribution to the university, and its
297 community (72). As a result of this implementation, the number of promotion applications increased by
298 50% and the number of women promoted has also increased.

299

300 The University of Glasgow's academic promotion criteria explicitly rewards researchers for participation
301 in peer review and other related activities (e.g., journal editorship) (73,74). In order for this to occur, it is

302 necessary to have organizations that can provide reviewers with a permanent identifier (a Digital Object
303 Identifier (DOI)) for journals that publish Open Reviews (75) that can be included in a researcher's CV or
304 which can aggregate completed peer reviews (76). Such policies might also help promote more
305 meaningful involvement in training in peer review (76). The University of Exeter, UK, has developed
306 'Exeter Academic', a hub to help their researchers navigate career progression (77). Leadership and
307 citizenship are two (of five) major areas of focus. The former includes mentoring and the latter includes
308 avenues to disseminate research knowledge from the university's researchers.

309
310 The Finnish Advisory Board on Research Integrity (TENK) template for researcher CVs includes a broad
311 spectrum of contributions including mentoring and 'trust in society' (78). As a measure of mentorship,
312 Maastricht University, The Netherlands assesses the career progression of its PhD graduates (79). We
313 were unable to identify research institutions that reward researchers who have participated in training
314 courses on high-quality mentorship (20).

315
316 The Irish Health Research Board (HRB) has a knowledge exchange and dissemination grant program
317 providing existing HRB-funded researchers with an opportunity to seek supplementary funding for
318 exchange and dissemination activities that can accelerate and maximize the potential translation and
319 impact of the research findings, and learning gained, on policy or practice and health outcomes (80). A
320 similar scheme exists through the Canadian Institutes of Health Research (81) and the NC3Rs Skills and
321 Knowledge Transfer grants (82) and their Crack IT open innovation platform (83).

322
323 Wellcome's grant forms limit the number of publications applicants can submit and explicitly invite
324 applicants to detail other achievements. This is combined with explicit guidance for panel members
325 reminding them of the importance of taking a broad view when assessing individuals (84).

326

327 **Discussion**

328 The HKPs focus on promoting assessment practices that strengthen research integrity by deliberately
329 concentrating primarily on what research institutions can do to modify the criteria used by PTCs for
330 career assessments. The five principles we formulated are aimed at how research institutions should
331 incentivize, reward and assess individual researchers within their respective organization. The HKPs do
332 not address gender and other forms of diversity, inclusiveness, and other related issues. These themes
333 require an assessment of a group of researchers (e.g., research institution) when making decisions about

334 funding allocations or human resources policies. Individual researchers are obviously not in a position to
335 change their gender, equity or diversity. Furthermore, these issues concern the social justice and
336 societal relevance of research rather than research integrity.

337

338 *Dissemination*

339 The World Conferences on Research Integrity (WCRI) Foundation (85) and the REduce research Waste
340 And Review Diligence (REWARD) Alliance (86) will make the HKPs available on their websites. This
341 'home' will include the principles, the signatories, infographics, translations into several languages
342 (ongoing), future implementation plans (ongoing), and crucially, a place to highlight those who have
343 endorsed the HKPs. Beyond journal publication, we are developing other synergistic dissemination
344 routes.

345

346 *Endorsement and Uptake*

347 Research institutions are key to the HKPs. They are the home of current and future researchers, where
348 promotion and tenure assessments are carried out. To help facilitate HKPs 'on the ground', local key
349 opinion leaders, and their endorsement, should be included in any plan. The HKPs have been recognized
350 by the Governing Board of the WCRI Foundation and the Steering Committee of the REWARD Alliance.
351 We invite academic institutions, funders, other groups and individuals to do likewise on the WCRI
352 Foundation's [website](#).

353

354 We are inviting individuals and organizations to deliver brief (2-3 minutes) YouTube testimonials as to
355 how they have implemented the HKPs (categorized by stakeholder group) and we will provide a link to
356 these videos on the WCRI Foundation website. This approach can serve as a pragmatic way for
357 individuals and organizations to show how they are endorsing and using the HKPs and as a nudge to
358 others to do likewise.

359

360 To implement some of these principles is likely straightforward although this might not be the case for
361 all principles. To do so requires more understanding of the complexities of today's research
362 environment, such as the availability of institutional infrastructure, whether current CV formats are
363 optimal to collect best practices, enabling transparency about career assessment, and considering closer
364 alignment with policies of funders.

365

366 We would like to evaluate our approach and develop tool kits for those interested in ways to implement
367 the five principles. We will work with signatories to take this forward. We see the HKPs as an important
368 step along the way to improving research integrity and we encourage an ongoing dialogue to support
369 implementation of these important principles.

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372 earlier versions of the document and actively participated in the focus group sessions during the
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374

375

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509

510 Figure¹: Robust, rigorous and transparent practice and impact

511

Research stage	Potential measures of rigorous research practice	Importance to research quality
Question	Knowledge synthesis Priority-setting exercise; stakeholder(s) engagement;	Useful and relevant research that builds on previous research
Design	Open protocols; (Pre)registration Reuse of protocol by others	Reduces publication bias and other reporting biases; Enhances reproducibility
Conduct	Quality assurance of data; Data sharing; sharing materials Reuse of data/materials by others	Allows data aggregation, data reuse, and transparency
Analysis	Analytical code sharing	Enhances reproducibility
Report	Transparency; open access; Use of reporting guidelines	Enhances openness and accessibility
Dissemination	Impact on research (including altmetrics; citations) Impact on practice/society	Focuses on outcomes & impact of research

512

513 ¹Items in black are measures of responsible research practice; items in red are measures of use by

514 others