

‘Prospecting responsible technology paths: Management options for an appropriate societal embedding of medical neuroimaging’

Marlous E. Arentshorst^{1,*}, Tjard de Cock Buning¹, Wouter P. C. Boon²
and Jacqueline E. W. Broerse¹

¹*Athena Institute for Research on Innovation and Communication in Health and Life Sciences, VU University Amsterdam, De Boelelaan 1085, 1081 HV Amsterdam, The Netherlands.*

²*Copernicus Institute of Sustainable Development, Utrecht University, Heidelberglaan 2, 3584 CS Utrecht, The Netherlands.*

**Corresponding author. Email: m.e.arentshorst@vu.nl.*

Responsible innovation implies an alignment of what developers and societal actors perceive to be the problems and purposes of new technologies. With this, the challenge is to prospectively identify those potential concerns and (systemic) barriers that might hamper the development and embedding of innovation. We address this challenge by contextualising different visions of medical neuroimaging, which we identified via interviews and focus groups. We show that different visions result in different desirable technology paths, each with specific concerns and barriers. Concerns include medicalisation and the burden of knowing a predisposition. Barriers comprise: scientific unknowns, technical impossibilities, disciplinary boundaries, and the focus on disease categories and cure in research and health practice. Proposed strategies to overcome the barriers include: different research incentives, training of scientists and health professionals, and developing person-centred health centres. We conclude with implications for the responsible management of medical neuroimaging, in which shared visions and mutual learning are key elements.

Keywords: responsible research and innovation; neuroimaging; emerging technologies; constructive technology assessment; vision assessment; technological paths.

1. Introduction

During the emergence of technological novelties, socio-institutional embedding and the technology itself co-evolve (Nelson and Winter 1982; Bijker et al. 1987). The development of an innovation is therefore dependent on interrelated dynamics and mechanisms, such as: the articulation of demands, networks and technologies. In this regard, it is important to take societal demands and concerns into account, as the failure of innovations such as genetically modified foods and crops in Europe and the

subsoil CO₂ storage in the Netherlands signified (Chilvers and Macnaghten 2011; Economic and Social Research Council 1999).

Taking these various forces into account during technology development should lead to innovations which target the creation of societal benefits, including economic growth, and also the management of the negative side effects of innovations on society and the natural environment. This perspective on technology development and innovation processes is increasingly referred to as

responsible innovation or responsible research and innovation (RRI). Von Schomberg (2011: 9) defines RRI as:

... a transparent, interactive process by which societal actors and innovators become mutually responsive to each other with a view to the (ethical) acceptability, sustainability and societal desirability of the innovation process and its marketable products (in order to allow a proper embedding of scientific and technological advances in our society).

Stilgoe et al (2013: 1570) take a broader perspective by indicating that it:

... means taking care of the future through collective stewardship of science and innovation in the present.

Managing innovation trajectories to facilitate an appropriate societal embedding is not a new phenomenon and diverse approaches incorporate RRI-related concepts, such as better foresight, more responsive and adaptive governance and public engagement, aiming to manage innovation process and open them to societal influence (Owen and Goldberg 2010). Constructive technology assessment (CTA) is an example of such an approach and aims to learn about the potential positive and negative impacts posed by a wide range of actors (Rip et al. 1995). CTA has a long history of aiming for technologies that connect better with societal practices, and has already been operationalised and implemented in practice since the late 1980s (Broerse 1998; Broerse et al. 2009; Rip et al. 1995; Roelofsen 2011; van Merkerk 2007). In this paper the focus is on CTA as an approach to RRI (see Section 3).

Applying approaches to RRI in an early phase of innovation development is expected to lead to an improved translation of the innovation and to facilitate its embedding in society (Wilsdon and Willis 2004; Rogers-Hayden and Pidgeon 2007; Roelofsen 2011). Research on CTA indicates that approaches aiming at an optimum balance between the desirable positive and undesirable negative impacts of innovations can add value to emerging and ongoing research and to reflection on research agendas (Rip 2009, Roelofsen 2011). However, the subsequent responses of actors are generally disappointing (Rip 2009; Roelofsen 2011; Schuurbiers and Fisher 2009). Although actors come to new insights and are stimulated to establish new spin-offs and adjust their research agenda, realisation of these intentions usually fails. One reason offered in evaluations of these approaches is that the intentions do not comply with the dominant culture (thinking), structure (organising) and practice (doing) that form the socio-technical regime (Geels 2004; Roelofsen 2011). Changing the dominant regime is complex, given its resilience to change. For example, Kloet et al. (2013) showed how the activities of actors in a multi-actor, innovation-centred consortium were constrained by the formal and informal rules, regulations and procedures of the socio-technical

regime, barriers that hinder the development and implementation of desirable applications can be identified, explained and, potentially, managed.

To deal with the barriers imposed by the incumbent regime and to contextualise the ideas and visions around an emerging technology, RRI implies the challenge of prospectively identifying potential concerns and (systemic) barriers that might hamper the development and embedding of emerging science and technology. Such an RRI process might lead to innovations in the direction that the relevant actors perceive to be desirable. In the light of the need to better understand how emerging technologies like medical neuroimaging innovations can be responsibly positioned *vis-à-vis* the incumbent regime, we took the challenge of prospectively identifying potential concerns and (systemic) barriers that might hamper development and embedding by operationalising CTA as an approach to RRI.

This paper focuses on neuroimaging innovations in the Dutch clinical context as an example of an emerging technology with potential benefits and negative effects. Innovations in neuroimaging make it possible to visualise and study the function, connectivity, activity and biochemistry of the brain as an intact structure (e.g. functional magnetic resonance imaging, positron emission tomography, electro-encephalograms and magneto-encephalography). To date, these technologies have contributed to insights into neural processes associated with three major types of disorders: psychiatric (Malhi and Lagopoulos 2007), behavioural (Dickstein et al. 2006), and degenerative (Rosas et al. 2004) brain disorders. Moreover, neuroimaging technologies have contributed to improved diagnosis and therapies for some of these disorders. They are primarily embedded in secondary care (i.e. hospitals), and applied to diagnose brain disorders and, to a lesser extent, assist in the treatment of some of these disorders. Technological advances, such as increased spatial and temporal resolution and improved options for data analysis, are expected to result in more detailed views of specific regions of the brain. This should lead to an increased understanding of both the brain and the origin and development of brain disorders. These technological advances and resulting knowledge are expected to lead to the development of improved diagnosis and treatment options and to contribute to novel options for prevention (Ewers et al. 2011; Szymanski et al. 2010; Willmann et al. 2008). Negative effects include the growing knowledge of the brain that may further extend the boundaries defining illness or redefining problems as medical conditions (medicalisation), and thereby increase the demand for medical services. People who do not display any symptoms may not want to know that they have a subclinical brain disorder, especially in the absence of an effective treatment (Fuchs 2006; Glannon 2006; Illes and Racine 2005). The positive and negative impacts of neuroimaging are uncertain in the current early phase of development.

In this paper we address the challenge of prospectively identifying concerns and (systemic) barriers that might hamper the development and embedding of medical neuroimaging by contextualising identified visions of actors related to these developments in the Netherlands and explore how these visions are embedded in the norms and values of the broader innovation community. The aim is to deduce those factors, mechanisms and dynamics which might become barriers during the development of neuroimaging innovations, thus adding to the understanding of how to formulate strategies to manage medical neuroimaging in a responsible way.

2. Towards prospective responsible technology paths

The early phases of innovation development are characterised by uncertainty regarding which developments will be realised, what knowledge will be generated, what artefacts are to be developed, and what societal impacts these might have. In this phase, technology developers may hold different visions (or beliefs) about ‘what is feasible or at least worth attempting’ (Nelson and Winter 1982: 258). This leads them to follow different paths to develop innovations (Garud and Rappa 1994). During development, specific technological competencies, such as the form and function of the artefact, are articulated. Technology developers within a group share perceptions of an attainable future and shape the future of technical artefacts with their ideas, through which a script of expected user behaviour is materialised in the artefacts (Akrich 1992). These perceptions and scripts function as a ‘language’ that guides actions in the concrete practices of technology development and are called ‘guiding visions’ (Grin and Grunwald 2000).

Guiding visions are not developed in isolation: the technology becomes institutionalised in a community of developers (Garud and Rappa 1994). Technology developers in the same group, who share certain guiding visions, also share the same rules, routines, and structures for evaluating the technology, resulting in a specific technology path (Garud and Rappa 1994) or technological trajectory (Dosi 1982). Technology developers tend to commit themselves to this technology path, becoming path dependent. Data consistent with the practices of the group are perceived as information and data inconsistent with the practices of the group are sometimes ignored or perceived as noise (Geels 2004). This is not a linear process, as has been pointed out by Garud and Rappa (1994). It is a process in which there is a reciprocal interaction between the beliefs or guiding visions of a technology developer (‘what is possible’), the technical artefacts they create (form and script or functional use of technology) and the evaluation routines they promote (testing routines and normative values).

Garud and Karnøe (2001) introduced the notion of ‘path creation’, which connects path dependence with studying alternative routes for technology development. The process of path creation includes deliberate or mindful reflection on possibilities for diverting from the existing path. The notion of path creation highlights that the actions of actors have consequences for the paths that are being made. In this way, actors influence path creation in a real-time manner. Thus, understanding the evolution of technology could provide options to manage an innovation in the early phases of its development.

When studying these prospective technology paths, the three interrelated elements of Garud and Rappa’s socio-cognitive model of technology evolution also play a role. In this context, visions about the technology serve as a proxy for the beliefs about potential future artefacts. These visions explicitly leave room for reflexivity, thereby extending an actor’s scope and evaluation criteria. The function of visions is to:

... deliver orientation for present acting and deciding. (Grin and Grunwald 2000: 179)

In other words, long-term considerations of future expectations are used to orient present actions, including those that influence technology development. Understanding these long-term orientations offers opportunities to identify emerging technological paths that are deployed by innovation developers. By identifying the visions of other relevant actors, it also offers possibilities to divert from these paths or to create alternative paths. In turn, the identification and combination of the individual visions of relevant actors offer opportunities to construct a more balanced, shared vision (Grin and Grunwald 2000), which serves as a signpost for a more responsible technology path.

Technology paths are also influenced by external factors (Geels 2004) such as: an increase in the number of chronically ill patients, the economic crisis and innovative initiatives. A system perspective provides an understanding of the mechanisms and dynamics of the socio-technical regime and facilitates an analysis of potential systemic barriers hampering the development and embedding of innovation (Kloet et al. 2013). This provides an understanding of whether the activities of actors are indeed imposed by the barriers of the incumbent regime, and if the strategies proposed to overcome the barriers take into account the dominant structures and practices of the system.

3. Method

3.1 Approach

To operationalise CTA as an approach to RRI, we use the interactive learning and action (ILA) model (Broerse and Bunders 2000), combined with vision assessment (Grin and Grunwald 2000). Roelofsen et al. (2008) showed that combining these two approaches results in an approach

which is suitable for analysis of, and intervention with, emerging technologies in the field of ecogenomics. We build upon these insights and apply this approach combined with a system perspective (Geels 2004) to assess medical neuroimaging technologies. The approach is characterised by an emergent and flexible design. Key features of the ILA model (Broerse and Bunders 2000) are: encouragement of active participation of all relevant actors, early in the process and on an equal footing; explicit acknowledgment of experiential knowledge; development of shared visions; knowledge creation through mutual learning (via dialogue); enhancement of trust relationships; coalition building; and independent and competent process facilitation. With vision assessment, shared future visions can be shaped that guide the directions of technology development (Grin and Grunwald 2000). Elements central to the identification and construction of visions (Grin and Grunwald 2000; Roelofsen 2011) include:

- *Problem definition*: different visions can entail various problem definitions and ways to assess solutions. Assessing the assumptions underlying a problem definition uncovers values and norms of how actors look upon reality, perceive facts and define the problem.
- *Challenges and purposes to be fulfilled*: This element concerns the challenges and purposes to be fulfilled, resulting from the specific practice in which the actors participate. The problem definition contextually vindicates the challenges and purposes to be fulfilled.
- *Relevant contextual aspects*: This element explores the relation between the technical artefact and contextual aspects. Examples include: the context in which the artefact will be used, how and by whom (e.g. conditions under which the technical artefact may contribute to solving a problem), who will benefit and who will possibly experience disadvantages. These elements also include factors that may hamper the realisation of the envisaged technical artefacts, namely barriers that need to be overcome.
- *Basic features of the desirable state*: This element refers to the basic assumptions around which visions develop: the preferred state of affairs the vision entails and ideas about what the world should look like.

This combined approach gave us the opportunity to identify those visions of neuroimaging from different relevant actors which related directly to the aims of our research, as described above.

3.2 Research design: Identification of neuroimaging visions

To identify the vision of neuroimaging from different relevant actors (see Box 1 for an overview), we started with the identification of future neuroimaging technology paths and potential resulting artefacts. We made an

Box 1. Actor field of medical neuroimaging We distinguish the following different actor groups which have their own structure, culture and practice and share structures with other groups forming together the wider societal health system:

- *Scientists*: actors who work with neuroimaging technologies or knowledge resulting from neuroimaging applications in a research setting.
- *Industrial producers*: actors who produce neuroimaging technologies. These actors may be concerned with technical standards and functional requirements.
- *(Potential) future users*
 - *Receivers*: actors who undergo neuroimaging (e.g. patients).
 - *Apppliers*: actors who apply neuroimaging in clinical practice or use the knowledge resulting from these technologies (e.g. health professionals). Within this group we distinguish the following health professionals based on the current organisation of the health system and differences in structures and practice on a more detailed level:
 - professionals working in primary care
 - professionals working in secondary care
 - professionals working in the field of somatic disorders
 - professionals working in the field of mental disorders
 - *Host institutions*: actors of neuroimaging companies and institutions in which neuroimaging equipment is located, including hospitals and private imaging institutes, who deal with liability and how to apply these technologies.
- *Policy-makers*: actors who deal with rules concerning administrative regulations and procedures which structure the health system. For example, regulations regarding the application of technologies, safety standards, and reimbursement.
- *Citizens*: actors who might use or are affected by neuroimaging in the future, but are not part of the health system. In contrast to the actors described above, the perceptions of citizens are based on a personal perspective rather than a professional one. Their knowledge can be considered as ‘contributory expertise’ (Collins and Evans 2002) and their desires, demands and concerns should also be taken into account at an early phase of innovation R&D in order to maximise the potential benefits of innovations for future users.

inventory of guiding visions from a neuroimaging developer's perspective, including scientists and industrial producers, because the beliefs and ideas of these developers currently shape the future directions of neuroimaging (Akrich 1992; Garud and Rappa 1994; Grin and Grunwald 2000; Roelofsen et al. 2010). Therefore we conducted and analysed semi-structured interviews ($n = 17$) and four focus groups with industrial producers, technology developers and scientists ($n = 19$) (details in Arentshorst et al. 2014a). We selected respondents who had specific expertise and experience as an individual representing an actor group. Throughout the process we challenged the interviewees and focus group participants to propose other relevant actors who were expected to be affected (positively or negatively) by neuroimaging. These actors were taken as a starting point for the next phase of the research. This 'snowball exercise' for recruiting respondents was then applied to identify and consult other actors and ended when referrals did not result in new suggestions. As a next phase of our research we identified and constructed the visions of potential relevant societal actors. For this purpose we interviewed five policy-makers, eight health professionals and three representatives of patients (details in Arentshorst et al. 2014b). In addition, six focus groups with Dutch citizens ($n = 46$) were organised to analyse public perceptions of medical neuroimaging (details in Arentshorst et al. 2014c). Next, we organised a multi-stakeholder dialogue meeting with scientists, industrial producers, policy-makers and health professionals ($n = 17$) to raise awareness of the different visions of neuroimaging and to validate and discuss our results. Questionnaires were then used to identify actions undertaken as result of the dialogue meeting.

As a consequence of the early phase of neuroimaging development, we consulted those actors who are most interested in new scientific and technological developments. These actors were capable of disembedding from existing structures (i.e. mindful deviation) (Garud and Karnøe 2001). Therefore they might be classified as pioneers, and it is likely that the views presented in this paper do not correspond with the views of other actors in the same field. However, pioneers, in contrast to their fellow regime actors, are those most likely to take up the challenge of establishing the changes they perceive to be necessary in the health system when options to do so become available. Thus, they might influence the future views and practice of actors in the same field and mobilize a collective, despite resistance and inertia (Garud and Karnøe 2001). They therefore seemed to be the best selection for this study. Although our study might not encompass all potential visions, we did obtain saturation with regard to articulated desirable artefacts and barriers, meaning that we only stopped interviewing different actors when we received no new information. The barriers related to the three different visions of neuroimaging have been validated by the respondents in our study as important elements in managing, prospectively,

neuroimaging developments towards more responsible artefacts, including their responsible societal embedding.

Interviews, focus groups and the dialogue meeting were audio-recorded and interviews and focus groups were transcribed verbatim for further analysis. Summaries of the interviews and focus groups were returned to the respondents with the request to check whether the summary was complete and interpretations made were correct. The identities of the respondents were anonymised by replacing their names with unique research codes. The data was then analysed using qualitative data analysis software (ATLAS.ti). In this analysis we focused on the identification of elements regarded as important in vision assessment: problem definition, challenge and purposes to be fulfilled, relevant contextual aspects and basic features of the desirable state. New sub-elements were noted as they became apparent in the data (Grin and Grunwald 2000; Roelofsen et al. 2008).

To gain an understanding of the challenges and barriers that might become obstacles when neuroimaging is further developed, we made use of the multi-level perspective (MLP) (Geels 2004; Loorbach 2007). The MLP is a heuristic tool to analyse the dynamics of socio-technical changes. Three interrelated analytical levels are distinguished: the landscape level (the exogenous and slow-changing landscape); the regime level (communities of interacting groups with dominant cultures, structures and practices); and the niche level (small-scale innovative initiatives). Placing our results in this framework allowed a contextualised analysis of the concerns and (systemic) barriers articulated from a system perspective. In Section 4 we describe identified concerns and barriers that might become obstacles when neuroimaging is further developed, followed by strategies described by the respondents as to how to manage these. We integrate the results of previous phases of our research (see Section 3.2) with this, and as a result make a distinction between the visions of: neuroimaging developers (comprising scientists and industrial producers), societal actors (comprising health professionals, policy-makers and patient representatives) and citizens.

4. Concerns and barriers that might complicate or hamper neuroimaging development and embedding

From a developer's perspective, desirable neuroimaging applications focus on new and improved options for the prevention, diagnosis and treatment of brain disorders (details in Arentshorst et al. 2014a). All the health professionals, policy-makers, patient representatives and citizens who were consulted also envisaged the formulated neuroimaging options from a developer's perspective, as being conditionally desirable in themselves (details in Arentshorst et al. 2014b; 2014c). But what about the concerns and barriers related to the envisioned desirable embedding of these neuroimaging

applications in practice? We now contextualise the identified visions of respondents and explore concerns and factors, mechanisms and dynamics which might become barriers during the development of neuroimaging innovations.

4.1 General concerns and undesirable neuroimaging paths

All the respondents raised concerns related to the enhancement of 'healthy' people, which potentially results in questioning what is normal and healthy. With this, they expressed a fear of the medicalisation of relatively healthy people. Furthermore, some citizens, patient representatives, health professionals and policy-makers expressed concerns with regard to the burden of frequent monitoring with respect to time and the challenge of confronting the finiteness of life. In addition, some citizens and patient representatives articulated concerns regarding the potential harmfulness of frequent neuroimaging use and a shift from testing medication on animals towards the acceptance of humans as test objects (e.g. neuroimaging possibilities to monitor the effectiveness of medication in the brain). Furthermore, all respondents expressed concerns regarding the potentially negative social and economic implications that an early diagnosis or indication of predisposition can cause. This could result in unethical, and therefore undesirable, use of neuroimaging artefacts, for example stigmatisation and discrimination caused by the use and abuse of neuroimaging data by commercial parties, such as insurance companies and mortgage lenders. Many citizens also found the use of neuroimaging to search for additional disorders, life styles and sexual preferences undesirable. Additionally, they expressed their fear of the reduction of a person to a mere 'image', excluding the experiences of health professionals, patients and nurture factors from the diagnostic and treatment process.

Concerning the use of neuroimaging to determine a predisposition, the uncertainty the predisposition represents, in terms of the chances of developing a disorder, and the burden of knowing were articulated as disadvantages of this artefact by all respondents, especially when no treatment is available. All neuroimaging developers, representatives of patients, secondary care professionals, and some of the citizens and policy-makers consulted, were concerned that large-scale preventive neuroimaging use might be expensive in terms of personnel and follow-up action and thus jeopardise the overall affordability and accessibility of health care.

Some health professionals, policy-makers, patient representatives and citizens expressed concerns with regard to the rules and regulations. On the one hand, they argued that rules and regulations often develop slowly, potentially resulting in applications being applied too early in ways that might turn out to be risky or unethical. For example, the Dutch law states that citizens with a genetic burden cannot be excluded from health insurance. However, it is

still possible to reward a symptomless group of people, which can lead to a financial disadvantage for those with a genetic burden or chronic disorder. As neuroimaging might give more insight into who has a predisposition or a disorder at an early stage, some of them expect that health insurance companies will increasingly explore these kinds of 'rewarding' options. They argue that the law should be adjusted before neuroimaging applications make this option possible. On the other hand, some health professionals and neuroimaging developers perceived that rules and regulations hamper the full exploitation of neuroimaging possibilities. An example is the European directive on the 'minimum health and safety requirements regarding the exposure of workers to the risks arising from physical agents (electromagnetic fields)'.¹ This directive, which is not yet implemented, addresses the protection of workers exposed to electromagnetic fields by setting a limit to the exposure to electromagnetic fields. However, the proposed maximum exposure for people working with MRI techniques is so low that making a MRI scan might no longer be possible.

The majority of the developers who were consulted (both scientists and industrial producers) were concerned about the management of expectations around neuroimaging innovations, which they perceived as a potential barrier. Some of the health professionals, policy-makers and patient representatives who were consulted were also concerned about the insufficient and/or exaggerated communication of the possibilities of neuroimaging to the general public. For example, claiming that many or all disorders and behaviours are located in the brain, which can be 'fixed' when there is something 'wrong', is an oversimplification. The communication of such a message to the general public is expected to result in inevitable disillusionment. The more science gives society the idea that 'they' know what is wrong and how it can be 'fixed', the more this will result in demands from society that science has to solve its problems. Neuroimaging developers considered misconceptions of neuroimaging by the general public to be a barrier that could lead, over time, to disillusionment and the rejection of neuroimaging technologies. These concerns are summarized in Table 1.

4.2 From different desirable neuroimaging paths to different barriers to solve

Although all respondents envisaged that neuroimaging applications would realise new and improved options for prevention, diagnosis and treatment, the desirable contextual aspects of potential applications and the underlying basic features of the desirable state are perceived differently. In other words, the technical artefacts in themselves are considered to be desirable, but the embedding envisaged in practice differs. Using data from interviews and the focus groups, we identified and constructed three different visions of how actors relate neuroimaging to the health

Table 1. Concerns formulated and strategies to overcome them

Vision	Concern	Strategy	Result of strategy
1–3	<ul style="list-style-type: none"> ● Enhancement and medicalisation of ‘healthy’ people. Harmfulness and burden of frequent use of neuroimaging ● Negative social and economic implications ● Reduction of a person to a mere ‘image’ ● Burden of knowing predisposition ● Overall affordability and accessibility of health care when preventive neuroimaging applications are applied on a large scale ● Slow development and hampering aspects of rules and regulations ● Management of expectations 	<p>Setting boundaries and rules for use of neuroimaging. Government is considered to be prime mover here</p> <p>Information dissemination:</p> <ul style="list-style-type: none"> ● Provision of information to individuals/ patients (e.g. regarding available options, reliability of tests, potential damage and consequences etc.) ● Provision of ‘correct’ information by government and scientists to health professionals and general public about possibilities and limitations of neuroimaging 	<p>Reduction of concerns</p> <p>Appropriate societal embedding of neuroimaging applications</p>

system. Neuroimaging technologies are envisioned in: current health care practice, personalised health care, and person-centred health centres (see Box 2, details in Arentshorst et al, 2014b).

In Vision 1, the technical optimisation of preventive, diagnostic and treatment tools is considered to optimise the structures and practices of the health system and hence contribute to a better health system in general. In addition to this technical optimisation, Visions 2 and 3 imply structural changes in the structures and practice of the health system and, through this, a better health system in general. In other words, the challenges and purposes to be fulfilled, the contextual aspects, and the underlying basic features of the desirable state differ, resulting in different visions for neuroimaging. In an earlier phase of our research we concluded that our results indicate that the visions of neuroimaging are not exclusively related to a specific group of actors, such as policy-makers or primary care professionals. We observed that the position that a respondent has in the health system and the vision he/she has of the ideal health system (which are interrelated) drive the vision of neuroimaging that an actor holds. As a result, desirable neuroimaging applications are envisaged in a way that they maintain or increase the position of a respondent, and suit his/her vision of the ideal health system and/or contribute to establishing this ideal health system, resulting in different desirable technical artefacts (e.g. a large device in a hospital versus a mobile device in a health centre) and related technology paths. Different technology paths to establish neuroimaging artefacts each have, in turn, specific barriers. The respondents who envisage neuroimaging in the current health care practice (Vision 1), formulated most of their challenges and barriers at a technological and knowledge level. Various respondents holding this vision also identified barriers intrinsic to the dominant science regime. Respondents who envision neuroimaging in personalised health care (Vision 2), and respondents

who envision neuroimaging in person-centred health centres (Vision 3), articulated, besides the aforementioned challenges and barriers, challenges and barriers related to the way in which health care is provided.

4.3 Knowledge and technological barriers

The identified barriers focus on scientific knowledge, which is not yet sufficient to establish desirable neuroimaging artefacts, and the high costs and time-consuming aspects of scientific research. The technological barriers, such as temporal and spatial resolution, the large size of neuroimaging technologies and differences between scan results (both differences in devices and methods and between individuals), also need to be improved in order to realise the desired neuroimaging artefacts. An example of a knowledge problem is the difficulty of differentiating between a healthy and a non-healthy brain at the individual level. In this case, differences are currently only visible when large groups are compared, or when one individual is followed over time, being its own control with respect to interventions with medicine or otherwise.

Some of the neuroimaging developers, policy-makers and health professionals who were consulted only saw these barriers from the perspective that they were obstacles that needed to be conquered, whereupon the neuroimaging artefact can be developed and subsequently embedded in the health system. As explained by one of the scientists:

We are constantly trying to improve our cognitive capacity, but there are probably ways to do that psycho-pharmacologically at some point. If we have the knowledge then we can improve memory and know what kind of pills we can administer to establish that.

However, other respondents argued that knowledge and technological barriers are the result of barriers intrinsic to the science regime. They warned against the focus on

Box 2. Three visions of how actors relate neuroimaging to the health system

Vision 1: Neuroimaging in current health care practice

In this vision, neuroimaging technologies are affordable and able to visualise brain disorders at a sub-disorder and/or individual level. Their appearance is not that different from current neuroimaging technologies (i.e. mostly large equipment located in hospitals). Applications make improved prevention, diagnosis and treatment of brain disorders possible and are perceived as *embedded in the current structures and practices* of the health system in order to optimise it.

Vision 2: Neuroimaging in personalised health care

In this vision, neuroimaging applications are affordable and able to visualise disorders at an individual level (i.e. person-centred applications). The purposes to be fulfilled by the neuroimaging applications correspond with Vision 1, i.e. improved prevention diagnosis and treatment of brain disorders. In this vision, neuroimaging applications are envisioned to be embedded in a person-centred health system. *Embedding is envisioned in primary and secondary care and/or in a new in-between setting* (e.g. ‘one and a half care’), as long as the structure and facilitates personalised care.

Vision 3: Neuroimaging in person-centred health centres

In this vision, neuroimaging applications are affordable, compact, mobile and able to visualise disorders at an individual level. The neuroimaging applications are *primarily embedded at health centres, at primary care level*. The purposes to be fulfilled by the applications correspond with the previous two visions (i.e. individual prevention and (more) personalised diagnosis and treatment). In addition, respondents holding this vision have the purpose to apply collective prevention strategies (e.g. the screening of symptomless people).

a specific (part of a) disorder in the current science regime, resulting in the fragmentation of scientific research and subsequent tunnel vision by researchers. This structure hinders the exploration and study of relations between disease categories, resulting in parallel research activities,

which might be unnecessary when, for example, the underlying disease mechanisms are similar. Moreover, answers (knowledge) and options resulting from cross-fertilisation activities, which might be beneficial for both research activities and clinical practice, might be missed.

Furthermore, structures regarding scientific validity (i.e. rules regarding scientific evidence) are perceived to be a barrier. The current ‘golden standard’ hampers research into possible subtypes of a disorder because this implies working with smaller, more specific, groups for which the criteria to set up the groups are unknown. As a result, research into subtypes would imply a loss of statistical power and consequently more expensive research (cf. research into rare diseases and personalised medicine). As explained by one of the health professionals working in the field of mental disorders:

Even in the new DSM [Diagnostic and Statistical Manual of Mental Disorders] that distinction [between different types of depression] is still not made. I think there is a kind of dogmatism in the research that maintains itself. [...] People have to let go of certain ideas of doing research in a particular way and establish a new approach in which interactions between certain vulnerabilities, gene level, personality level and certain environmental variables have a central place. That is, of course, much more difficult because you need huge numbers of people. So I think in terms of costs, the research as it happens is a lot cheaper. If you break down those groups, subdivide them, then you lose evidence/statistical power. So then you need to have larger groups, which cost more money.

In addition, some of the developers and many of the health professionals, policy-makers and patient representatives who were consulted saw the disciplinary boundaries of the science system as a barrier. This barrier constrains the realisation of interdisciplinary research, which is considered to be necessary to advance the field. They perceived the scientific regime to be rigid with its own dynamics, within which many scientists strive for personal gain and status. This barrier is partly related to professional and financial structures, but it is also related to the reluctance of scientists to cooperate with other actors, both academics as well as non-academics. Paramedical professionals, for example, still face a huge struggle in becoming a partner in scientific research, because their research centres are connected to higher vocational rather than academic institutes. Many of the neuroimaging developers consulted argued that interdisciplinary research, although necessary, is expensive and time-consuming, which serves as a barrier to conducting this type of research. Moreover, interdisciplinary research is considered not to be ‘rewarding’. This is related to the lack of suitable journals that publish results from interdisciplinary research and the lower impact factors these journals have, which constrain career advancement.

4.4 Barriers intrinsic to the structures and practices of the health system

Additionally, respondents visioning neuroimaging in personalised health care (Vision 2) and in person-centred health centres (Vision 3), described barriers related to the structures and practices of the health system. These barriers arise from criticism of the current focus on disease categories in health care practice and focus primarily on the practice through which health care is provided. Respondents explained that many health professionals diagnose and treat diseases per category and hence focus on the disorder a patient has, rather than on the patient who has a disorder. According to them, such a focus results in the barrier of a limited vision of the illness and health of the patient in question and, as a consequence, in cases of co-morbidity, in the provision of contradictory cure and care options for the patient by different disciplinary experts. As explained by a representative of primary care professionals:

People often do not have a disorder, you deal with multi-morbidity [...] What matters is: what is wrong with this person? So not disease-specific and resulting conflicting advice, medication, etcetera. One should look at the whole person.

In addition, some health professionals, policy-makers and patient representatives described the barrier of patients feeling that they have no say in their diagnosis and treatment trajectory. This results, for example, in non-compliance and a passive attitude, because patients are not offered options to become responsible for their own health and recovery. Furthermore, the distinction that is still made by medical professionals, patients and society in general, between somatic and mental disorders was articulated as a barrier by many of the health professionals, policy-makers and patient representatives who were consulted. Somatic disorders (e.g. neurological disorders) are perceived as ‘real’ disorders and mental disorders (e.g. depression or anxiety disorders) are at least envisioned as ‘not that real’. According to these respondents, this results in a view of, respectively, patients who have a disorder and those who have complaints, and medical professionals who deserve more respect and status than others. As explained by a health professional working in secondary care in the field of mental disorders:

I think that the professionals have to communicate that more clearly [that a mental disorder is also a disease]. The problem is that many colleagues, and somatic professionals in particular, envision to some extent that psychiatry is also a bit of a trifle thing.

Furthermore, respondents holding the vision of neuroimaging in person-centred health centres (Vision 3), articulated the barrier that the focus of the health system is too much on cure, hampering the prevention of disorders and resulting in a distinction between primary and secondary care. As explained by one of the policy-makers:

... from care and illness to health and behaviour. First things first: Earlier! Quicker! Better! Then you need diagnostics and also screening.

The barriers and strategies formulated to overcome them are summarized in Table 2.

4.5 Management of concerns

According to the majority of the respondents, most concerns can be managed by setting boundaries and rules for the use of neuroimaging. For example, they argued that neuroimaging can be limited to those brain disorders for which a predisposition can be identified and restrictions can be imposed on the use of data resulting from neuroimaging outside the health system. Furthermore, according to the health professionals, policy-makers, patient representatives and citizens who were consulted, the concerns regarding the burden of knowing an early diagnosis and predisposition might be tackled by: setting rules and boundaries relating to when and how these neuroimaging artefacts can be used; and providing individuals with information with which to make a fully informed decision, such as: the available options, the reliability of the tests, the potential damage and consequences.

All the neuroimaging developers who were consulted, indicated that worries about unrealistic societal expectations can be reduced by providing the general public, including the government and health professionals, with correct information about the possibilities and limitations of neuroimaging. They thought that the government had a significant role to play in this process, but also envisaged a role for the scientific community to correct ‘false’ stories and to provide information to citizens and end-users. The majority of the citizens who were consulted indicated that more information regarding the possibilities and limitations of neuroimaging, and medical technologies in general, would be appreciated. Several health professionals, policy-makers and patient representatives mentioned that information regarding the anticipated possibilities and meaning of neuroimaging artefacts (e.g. ‘what does it measure’) might provide a clearer view of the possibilities and impossibilities of neuroimaging artefacts and prevent unrealistic expectations of medical professionals, patients and citizens. According to these respondents, the designated point of departure for this is the science regime.

4.6 Management of barriers

4.6.1 Knowledge and technological barriers. According to all respondents, the science regime will rise to the challenge to produce the required knowledge and will, together with industrial producers, establish ways to technologically improve neuroimaging technologies in order to eventually produce desirable neuroimaging

Table 2. Barriers and strategies to overcome them

Vision	Barrier	Strategy	Result of strategy
Knowledge and technological barriers			
1–3	<ul style="list-style-type: none"> ● Scientific unknowns ● Technological impossibilities 	<ul style="list-style-type: none"> ● More funding ● Public–private partnerships 	<p>More research Neuroimaging technological breakthroughs</p>
1–3	<ul style="list-style-type: none"> ● Barriers intrinsic to science regime: <ul style="list-style-type: none"> ○ Focus on disorders ○ Disciplinary boundaries of science regime ○ Scientific validity 	<ul style="list-style-type: none"> ● Change incentive structures to favour research: <ul style="list-style-type: none"> ○ Interdisciplinary research as a pre-requisite for funding ○ Interdisciplinary journals with high impact factor ● Education of scientists ● Development of statistical tests for small populations in relation to health risks for that population 	<p>More person-centred, interdisciplinary structure and practice of research</p> <p>Scientific evidence based on n = 1</p>
Barriers intrinsic to structures and practices of health system			
2 and 3	<ul style="list-style-type: none"> ● Focus on disease categories: <ul style="list-style-type: none"> ○ Distinction between somatic and mental disorders ● Patients have no say in diagnosis and treatment 	<ul style="list-style-type: none"> ● Development of integrated cure and care plans and implementation in interdisciplinary teams ● Education of professionals to perform new and different tasks in person-centred context ● New categories of professionals to apply and interpret neuroimaging in person-centred context ● Implementation of shared-decision making 	<p>Person-centred, integrated health care</p>
3	<ul style="list-style-type: none"> ● Clinical focus on cure: <ul style="list-style-type: none"> ○ Distinction between prevention and cure, and care ○ Distinction between primary and secondary care 	<ul style="list-style-type: none"> ● New health institutions at primary care level that are person-centred and interdisciplinary ● Development and implementation of self-management options ● Include shift towards primary care and self-management in governmental policies 	<p>Health system with a focus on prevention Shift from secondary to primary care and self-management</p>
1–3	Barriers intrinsic to science regime and structures and practices of health system	<ul style="list-style-type: none"> ● Networking ● Mutual learning 	<p>Concerted effort to realise different structures and practices</p>

artefacts (cf. public–private partnerships). At the same time, the health professionals, policy-makers, patient representatives and citizens who were consulted, emphasised that it is important that the scientists and industrial producers become aware of perceived undesirable neuroimaging use and of concerns to promote the development of responsible innovations. In addition, according to the majority of the respondents, financial support is required to overcome scientific unknowns and technological barriers.

All respondents promoting a vision of neuroimaging in personalised health care (Vision 2), in person-centred health centres (Vision 3) and some respondents holding

the vision of neuroimaging in current health care practice (Vision 1), considered that changes are necessary in the science regime to overcome knowledge and technological barriers. These respondents perceived that the science regime needed to adopt a different, disease transcending, interdisciplinary structure and practice of research. According to them, professional and financial structures should be changed in such a way that an interdisciplinary structure and practice of research becomes the new ‘business-as-usual’. This includes the education of scientists to change their way of thinking and practice, and the incorporation of this new way of thinking in the educational programmes of universities. The government and

funding agencies are seen here as the designated point of departure, for example, by developing new funding strategies in which interdisciplinary research is a prerequisite for obtaining grants. Furthermore, many of them emphasised the importance of a change in structures with respect to 'scientific validity' and related 'rewarding' of interdisciplinary research. Structures, for example demands for statistical power, should change in order to make research into subtypes of disorders possible, rewarding and accepted as scientifically valid evidence. More investment should be made in the development and dissemination of statistical tests for small populations in relation to the health risks for such populations. Furthermore, the creation of interdisciplinary journals with high impact factors was a frequently articulated strategy, aimed at making interdisciplinary research more rewarding. They argued that this new culture, structure and practice would enable the research activities that are perceived to be necessary to move from large patient groups to the individual level, that is, studying the relations between disease categories, cross-fertilisation studies and research into possible subtypes of a disorder.

4.6.2 Barriers intrinsic to the health system.

According to respondents holding the vision of neuroimaging in personalised health care (Vision 2) or in person-centred health centres (Vision 3), overcoming the perceived barriers related to the health system requires a shift in structures and practices from a disease-oriented towards a person-centred health system. These respondents argue that, given this change, the advantages of neuroimaging possibilities can be fully exploited. They suggested that this can be accomplished by developing integrated cure and care plans and involving interdisciplinary teams, which have the consumer/client/patient at their heart. To this end, professionals need to be educated to perform new and different tasks in a person-centred context and hence change their professionals practice (i.e. behaviour) accordingly. New categories of professionals should be developed to apply and interpret neuroimaging in this context, and future patients should be encouraged to change their behaviour and to become responsible for their own health, through, for example, self-management options.

Respondents holding the vision of neuroimaging in person-centred health centres (Vision 3) argued that, besides a shift towards a person-centred health system, a shift towards primary care and self-management is necessary. For this, a range of primary care professionals should be placed together in interdisciplinary health centres at primary care level. Respondents holding this vision argue that primary care should be the location where integrated care, including diagnostics, will start and, when appropriate, actors (professionals) from secondary care will be consulted or treat the patient. Moreover, self-management options should be offered to both patients and citizens to

become responsible for their own health and, if necessary, for their recovery trajectory. The government is again perceived to be the designated point of departure by pursuing these changes in their policies.

Dialogue meetings in which actors from different disciplines and regimes meet and learn from each other in a learning environment, might contribute to developing more shared technology paths and applications, according to participants who attended our dialogue meeting. Participants in the dialogue meeting reported that the meeting resulted in greater awareness and exploration of potential barriers and new areas of innovation by discussing desirable and undesirable applications with actors outside their own practice. However, as indicated by the participants, actors from different disciplines and regimes do not normally make the effort to have a dialogue with each other. As suggested by the participants, these dialogues require facilitation to both bring people together and to create a safe environment where mutual learning may take place, and such interactions need to be organised with some regularity.

5. Conclusions and discussion

5.1 Conclusions

In this study we explored concerns and barriers related to the development and embedding of neuroimaging in the Dutch health system from the points of view of different actors and from a system perspective. Our results show that all respondents have concerns related to the enhancement and medicalisation of 'healthy' people, the embedding of new preventive and diagnostic applications for which no therapeutic options yet exist, and the potential negative social and economic implications associated with preventive neuroimaging applications. Compared to neuroimaging developers, health professionals, policy-makers, patient representatives and citizens expressed more concerns about how neuroimaging may contribute to solving problems. This particularly applies to preventive neuroimaging use. According to many of the respondents, concerns can be managed when boundaries and rules are set for neuroimaging use and when sufficient and unbiased information is provided to those who undergo neuroimaging so that they can make a fully informed decision. With this they assume that when neuroimaging applications are well organised and there is proper information dissemination, concerns are 'managed' sufficiently. However, these strategies are traditional end-of-pipe policy tools of command and control with an underlying assumption that knowledge generates public acceptance of science and technology, and hence facilitates embedding of innovations (Druckman and Bolsen 2011; Nisbet and Goidel 2007). Previous research has shown that command and control measures and strategies to augment the knowledge of the general public result in neither a reduction of

societal concerns nor increased public acceptance (Chilvers and Macnaghten 2011; Hagendijk and Irwin 2006; Marris et al. 2001). Concerns related to the potential harmfulness of frequent use of neuroimaging and the technical ability to determine the ‘right’ diagnosis, should therefore not be dismissed as a lack of scientific knowledge and/or as emotional concerns that can subsequently be addressed by the dissemination of information. Instead, these concerns should be taken seriously, for example via the interaction of different actors in a dialogue, and be taken up in the technology development path.

This study shows which barriers need to be overcome in order to realise desirable neuroimaging applications. It demonstrates that different visions of neuroimaging result in different technology paths, which have their specific barriers. These barriers are cumulative and increase in number and complexity from Vision 1 to Vision 3. Respondents holding the vision of neuroimaging in the current health care practice (Vision 1) formulated most of their challenges and barriers on technological, knowledge and financial levels. Some of the developers, health professionals and policy-makers who were consulted formulated these barriers from the point of view that they are obstacles that need to be conquered, whereupon the neuroimaging artefact can be developed and subsequently be embedded in the health system in order to optimise it. For other respondents, these barriers are also related to the incumbent science regime. Respondents holding the vision of neuroimaging in personalised health care (Vision 2) and person-centred health centres (Vision 3) additionally articulated barriers and challenges related to the health system. These barriers are formulated from a point of view that structural changes in the health care practice are needed in order to appropriately embed neuroimaging applications and thus establish a ‘better’ health system. Indeed, many of the barriers are systemic in nature.

5.2 Towards an appropriate societal embedding of medical neuroimaging

We observed that the strategies articulated to overcome the barriers were formulated from within the professional practice of a respondent. They were envisaged to increase or maintain the position and/or status an actor has, while responsibility for the strategies was primarily handed to actors of other practices and regimes, preferably the government. In other words, actors protect and reinforce the rules of their regime (Geels 2004). The strategies are formulated from the actors’ perspective of the problem, without considering other actors’ perspectives. These strategies might therefore fail due to their focus on the individual’s own professional practice and their mismatch with the current structures and practices of the health system. In other words, although respondents identified barriers of a systemic nature, their strategies to overcome these do not take the resilient system into

account. On the other hand, there are potential destabilisers of the dominant regime. Landscape-level trends, such as aging which lead to more patients with brain disorders, and an increasing preference from citizens for having more control over one’s own health (Jones 2008), combined with overcoming barriers by, for example, making it financially and technically possible to shift towards person-centred approaches and/or primary care, are potential destabilisers of the dominant regime. The above findings indicate that actors who envisage neuroimaging in a health system with new structures and practices, might in this case start acting on the changes they perceive to be necessary. In that case, the incongruences in visions might be in conflict with each other.

In order to realise more responsible neuroimaging applications, actors need to make an effort to consider, understand and integrate the view of the other. To formulate strategies that take the resilience of the dominant system into account, the results of this research indicate that learning and understanding of the context from which barriers arise might result in strategies that are effective in overcoming the barriers (Regeer 2010). Understanding the different visions of actors, including perceived barriers and strategies to overcome these from a system perspective, offers opportunities to combine these visions constructively into a more balanced, shared and responsible vision (Grin and Grunwald 2000). Multi-actor dialogue meetings are perceived by some of the respondents as a strategy to this end. Becoming aware of the fact that one reinforces one’s own function and its practice, might result in discussions about what the future health system should look like, where neuroimaging is envisaged to be a part. In other words, underlying the arguments used to discuss strategies to overcome the barriers, there are assumptions regarding the basic features of a desirable health system and the functioning of the actors within this system.

Our research shows that a next step would be the development and implementation of an action plan through reflexive learning cycles of planning, action, observation, reflection and re-planning. This might shape technology paths towards more responsible paths and in turn, support the process of their embedding in society. What this step will look like in detail depends on the shared, desirable visions identified. It is important to note that such visions are neither a ‘final state’ nor a consensus goal, but more a guideline for responsible monitoring. After all, as emphasised in Section 2, innovations, artefacts and their socio-institutional context co-evolve during development, implementation, and use (Nelson and Winter 1982; Schot and Rip 1997).

We started from the conclusions of previous research that the ideas of actors as a result of a CTA process do not, in general, result in major changes to the innovation process (Grunwald 2011; Kloet 2011; Roelofsen 2011; van Merkerk 2007), probably because the socio-technical regimes are too dominant and too rigid to allow changes

(Kloet et al. 2013). Therefore, we analysed systemic barriers proactively throughout the process and attempted to raise awareness and facilitate the action of relevant actors with regard to the existence of different visions and related (systemic) barriers to facilitate the realisation of desirable neuroimaging applications. Whether this process results in long-term effects and what these effects are, cannot yet be assessed. However, experiences with multi-actor processes such as CTA, indicate that approaches to RRI should be viewed and designed as a continual learning process (Kloet 2011; Roelofsen 2011; van Merkerk 2007), in which the dimensions of anticipation, reflexivity, inclusion and responsiveness are integrated and institutionalised in and around the process of innovation development and embedding (Owen et al. 2012; Stilgoe et al. 2013). In this sense, the responsibility for and governance of, RRI process is regarded as a cooperative and distributive effort: the interactions during the learning process should lead to shared, guiding visions. Our results support this conclusion and suggest that processes aiming to use RRI in order to facilitate an appropriate embedding of the resulting artefacts, such as the iterative ILA model, could, and perhaps should, have a structural place in any emerging science and innovation which aims to produce societal benefits. This begs the foundational question: who is responsible for initialising and financing RRI processes? The majority of the respondents in our research envisage the government and funding agencies as the designated point of departure, as the science regime has no internal drive to change the existing focus on knowledge production in the absence of a central driver. In our view, a strong innovation research policy, demanding RRI designs for admissible research proposals, coaching the granted multi-disciplinary research teams and conditional mid-term evaluation on societal impact might be beneficial for wider circles of actors and society at large. By using such facilitation, a structural place for RRI processes in emerging science and technology (i.e. addressing related systemic barriers), might result in a cooperative and distributive effort for accountable financing with public money in public-private innovations.

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1. Directive 2004/40/EC, directive 2008/46/EC and directive 2012/11/EU see <www.ec.europa.eu> accessed 27 March 2014.

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