Holism’s fuller engagement with realities is an aspiration and ideal. In practice, there are many obstructions. These range from our use of language to our highly managed and industrialised culture. How does this happen? What are the consequences?
Prologue

Holism (and its lack) may be easier to recognise than define. It is more readily communicated and perceived by stories, rather than data or abstract formulations. This presents problems: holistic mindsets are now becoming harder to access and maintain, for our culture is now one that increasingly conceives and conveys in packages: food, fuel, news, entertainment, even thought are all likely to be coded, metered, monitored, measured or packed. This causes fewer problems when our encounters are with inanimate and less complex life-forms: the production and distribution of eggs or detergents cause fewer ethical and social conundræs than the industrialisation of complex welfare activities (though even our simpler activities eventually confront us with wider ecological – ultimately Gaian – consequences).

We thus have an insoluble handicap. It is always easier to think in parts than wholes: language, analytical thinking, our micro and macro economies … all tend to fragment our perceptions and activities: ‘this is this, and that is that’. In contrast, holism’s tenet of infinite and often hidden interconnectedness tends to erase boundaries and conflate territories: ‘this is that as well as this’. Such thinking largely eludes schemes, packaging, academia, economic analyses. Our use of language, too, struggles to convey any sense of holism without serious loss or distortion.

The following collage of notions illustrates, very partially, the extent of our difficulties and task. The notions themselves are presented without usual conventions of academic thoroughness or cohesion. The first two present the skeleton of the view: the briefer, later sections provide some illustrations and variations. Overall, they represent some unsystematised, though summative, personal reflections from one practitioner’s decades of working in human healthcare – a chimeric and often paradoxical world. Philosophical contention is ever-present. We are accelerating our mandates for factory-like language and procedures to service increasingly complex healthcare: human nature and predicaments remain considerably more ambiguous.
1. I’ve got a measurement – it must be a fact

The rise of data and the curse of scientism

‘Nothing vast enters the lives of mortals without a curse’

– Sophocles (c 496-405 BC)

There was life and technical success before computers, yet these are rapidly becoming harder to understand. Some examples: the manufacture of antibiotics, the D-Day Landings, Man on the Moon, Concorde – all of these were achieved with minuscule or no computer-power – things we could not manage now in our ‘progress’. We have become empowered but deskilled: in healthcare, as we shall see, these subtle discrepancies lead to grievous losses.

Before the widespread use of computers, the harvesting and collation of measurements – data – was manual, labour-intensive and therefore slow. It thus required much deliberation and discrimination and – relative to today – its volume was tiny and consequently much more manageable.

The electronic unshackling of these activities has freed them from the constraints of our individual capacities for engagement, assimilation or understanding: data has multiplied exponentially and is now pumped and piped at us like gas or water – public commodities.

Measurement, the blood-brother of data, has thus been conferred pre-eminent status in many humanly-complex activities. Numbers are the most easily digested ‘food’ for computers, and computers are now essential to the functioning of any public service. Existence of people and their activities must be continually monitored and broadcast in a form that can ensure their organisational recognition, management and survival. The virtual world now defines and commands the real: measure or perish. Once started, this is difficult to slow or stop.

So, our institutions are now electronically held together by computers, computers need data, data need statistics, statistics need measurements; ergo: measurement becomes the basic language and activity.
What does this mandatory measurement mean for healthcare? The consequences vary greatly with the type of activity. Sometimes the effect is facilitating and benign. For example, with activities that can be easily and directly measured, standardised and proceduralised: here the measurement culture can be applied with relative ease and evident benefit. Laboratory services, vaccinations and cataract extractions all serve as common examples. All have in common a clear, circumscribed physical basis, little variation in technique or human response and a high completion/success rate. In short, they can be easily humanly ‘mechanised’.

But much of healthcare does not offer this kind of simplicity for measurement, and then the effects often depart widely from the benign and facilitative. Measurements are at their most competent with physical objects or phenomena: a blood-count is far less problematic or contentious than a mood-rating scale. This is because attempts to assess, measure and code other people’s experiences must be derived from something else: self-reports, or other people’s perception being the commonest. All are subject to massive contention, contamination and compromise. What does this mean? Here are some personalised examples:

• Ms B is in dispute with the Department for Work and Pensions (DWP) over her Disability Living Allowance. Ms B claims severe symptoms and invalidity from Depression, but her invalidity seems invalid to the DWP assessor: he asks the GP, Dr F, for his opinion. Dr F, in a complex but rapid judgement, agrees with the assessor. This ‘objective’ assessment is stymied by Ms B’s continually high self-reported, but quantified, depression scores. ‘Something’s wrong’ they all say in different ways. ‘Only the wearer knows where the shoe pinches’ says an old adage, but when is this untrue? Who decides? How?

• Kenny is sixty-two years old, a single, lonely man, appeasing and self-deprecating in his manner. Harsh and neglectful parenting left him with impoverished self-esteem. A working lifetime as a road worker has riddled his lower body with degenerative arthritis. He left school at fourteen: his
intelligence exceeds his words. After several years of courteous wariness he is, with Dr F’s gentle encouragement, beginning to talk of his burden of fear, loneliness, shame and longing. The ancient story behind it is poignant and powerful. Kenny has great faith in Dr F, but continues anguished in his small and crumbling world. Dr F asks for help from NHS Psychological Services, to help Kenny occupy his limited life more positively. Kenny returns to Dr F a fortnight later, fearful, tearful and trembling. He nervously indicates the immediate cause of his distress: a tightly-stuffed, freshly opened envelope. It is from the Psychology Services. Dr F surveys several detailed questionnaires aiming to define diagnosis, severity, disability and numerous personal and demographic details. In addition are various bureaucratically prolix letters and documents explaining ‘The Service’, a Complaints Procedure and instructions for the Service User. No one has spoken to him.

In his frightened and faltering language Kenny conveys to Dr F his sense of bewildered humiliation and abject inadequacy: ‘I don’t understand all this, doctor … I just can’t do it … I just want to talk to someone – like I do with you, doctor.

The doctor remembers many years ago reading of Heisenberg, an early 20th century physicist. Heisenberg found that it was impossible to plot simultaneously the velocity and locus of an electron without changing these in an indeterminate way: the observation changed the reality. Dr F as a young man could not identify how this was relevant to his meagre knowledge of physics. Many years later he is seeing clearly how personal observation – when formulaic and non-bespoke – can adversely affect people he knows well.

• Philip is eighty-six and Dr F is visiting him at home, the week after his discharge from hospital. He had taken a first-ever overdose of his medication to end his life. An earlier than expected visit from his Carer had found him collapsed and vomiting.

Philip now looks tired and Dr F again senses immense melancholia beneath the mask of rigid discipline, of understatement. The doctor knows some of Philip’s recent trials and sorrows: his wife’s gruelling and fatal malignant illness,
followed rapidly by the sudden death of his beloved son, their only child. And then the increasing impoverishment of his own Parkinson’s Disease, a gathering bass-note.

Dr F had premonitioned Philip’s trapped but mute anguish and its possible tragic fruition, and had asked for help from his mental health services. Their (non) engagement proceeded by asking Philip to fill in detailed mood and anxiety questionnaires. These indicated mild, stable disturbance – measurements meriting merely a brief psychological care package from a Low Intensity (skills?) Worker, and a routine, templated report of all this, electronically conveyed to Dr F.

Dr F’s perception is discrepant. He shares his unease with Philip, whose intelligence and insight survive his ravages of grief: ‘I don’t like to tell anyone my troubles, doctor … I wasn’t brought up like that. I have my pride, you know … It’s different with you: I’ve known you years, and I don’t have to say much, for you to understand. But answer all those questions for a stranger? No.’

Dr F thinks of Philip’s formative years: a harsher, crueller, braver world of much greater trials, losses and endurance; a black-and-white world where contained and stoic fortitude was a social essential. Dr F understands this with few words, and Philip understands that he understands. But a questionnaire?

Yet Dr F now inhabits a professional world in thrall to designated experts who are keen to quickly code and quantify the distress of Philip and Kenny, as well as Dr F’s ministrations, and then to instruct them all. Dr F’s understanding can seem piecemeal, slow and never finished: features of the intersubjective, one-person-and-situation-at-a-time. By contrast the questionnaire has slick allure: its ‘objectivity’ may be specious, but it is quantifiable and can be given to all – a demotic science. Dr F is thinking of the distinction between the scientific and the scientistic: which is which? He is thinking, too, beyond his own professional end: who then will be speaking to the Philips of the world, and what kind of conversations will they be?

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2. I’ve got a word – it must be real
The trap of reification

‘In the Beginning was the Word’
– John, 1:1

Language does far more than merely ‘communicate’: words first contain, then command and control our experience, and then our influence of others. The implications of this for healthcare are subtle, powerful and rarely discussed. A brief linguistic analysis will help us understand these.

All words are there to package and convey a description of, or notion about, human experience. All ultimately come from our perceptions, then our constructions. The basic components of language are: adjectives, describing qualities of experience (what something is like); verbs, describing activity producing change (what something does); and nouns, which attempt to capture a more static state, a ‘something’, from which these other two emanate (what something ‘is’). For example, I describe a small vertical platform supported by four vertical supporting posts: someone comes and sits on it – it is a ‘chair’. Generally, we think such ‘real’ things endure and we attribute them by nouns: adjectives and verbs are more the flux of experience.

In our usual waking life this may present few problems, with nouns seamlessly providing apposite bridges and anchors for the rest of our sense-experience, and those of others. But potential dislocation is ever-present.

An example: I am at a friend’s table, eating an unfamiliar dish. I do not recognise the texture or flavour of the meat, although I enjoy both. I enquire what ‘it’ is.

#1. I am told: ‘It is lamb’. I am mellow with appreciation for my sensations, my friend and the cosmos.
#2. I am told: ‘It is Alginon, my ginger cat. She was very old and was
dying anyway’. What is in my mouth now triggers an explosion of nausea,
and retching. I jump up with disgust and mistrust. My friend and the
cosmos turned malign.

The ‘actual’ experience is transformed by the idea of the ‘real’ source-object
(lamb or cat). This noun now determines my subsequent experience and
action: the adjectives (pleasure or revulsion) and the verbs (sitting and
eating or jumping up and retching). All this happens despite my never
seeing the putative lamb or cat: they are abstractions rendered powerfully
‘real’ by the noun. Such is the power and gravitational force of nouns.

Nouns work with greatest clarity and efficiency when applied to physical
objects: the words ‘table’ or ‘television’ rarely cause problems except to a
foreigner, a lawyer or certain kinds of academic. We generally accept these
object-nouns as ‘real’. Elsewhere the use of nouns is more problematic and
more interesting: God, democracy or love may sound like (sacred) ‘things’,
yet are essentially variegated ideas. Innumerable stories from world or
domestic histories show how little clarity and consensus the nouns manage
here, yet how real they are to their believers.

There are striking analogies in healthcare. One working definition of
medical diagnosis is the organisation, then transformation of adjectives (a)
and verbs (v) into professionally conferred nouns (n) which then determine
explanation, therapeutic action and prediction for others. With afflictions
that are predominantly physical – ‘structural pathology’ or disease – we
can call this type of noun a ‘Substantial Diagnosis’. Here is a simple
example.

- Tommy is six. Last night he became listless (a), pushed away (v) his favourite
  supper, complained of soreness (a) in his throat and abdomen and then started
to shiver and vomit (v). Dr Y is now with Tommy and his mother. His job is to
  find and then confer the right organising noun, or diagnosis. When he sees
  Tommy’s much enlarged, reddened tonsils, flecked with creamy pus, he has the
  precise constellated noun: ‘Acute Pustular Tonsillitis’, though he thinks
‘Tonsillitis’ sufficient for his verbal communications. The formulation and conveyance of this word are beneficial for all: Dr Y knows what to do and what to expect, Tommy will almost certainly get better, Mother is comforted by this and the containing, reassuring clarity of this noun – the Substantial Diagnosis. For all, this process is helpful and uncontentious: the doctor’s knowing and naming the ‘thing’ of Tonsillitis is a cooperative and shared blessing. Importantly, Dr Y’s diagnosis also relieves the sufferers from having to search for their own explanation, meaning of, or influence on, events.

In other areas of healthcare this hegemonic use of nouns runs into many more difficulties. This is particularly so where the doctor is dealing with bodily dysfunction (functional dis-ease) in the absence of the evident structural changes of bodily disease. Equal difficulties are encountered with disorders of behaviour, appetite, mood or impulse (BAMI): the core of psychiatry and clinical psychology. The results here are more mixed: our medical-noun type diagnosis may sometimes bring evident clarity and relief to these physically non-fixated forms of distress, but often it will not. Then the professionally conferred noun – the diagnosis – is conveyed, but the benefits do not follow. In these situations the diagnosis may be ‘correct’, but clarification, relief or prediction remain poor. The doctor has – by convention – done his job, but none of the participants are gratified. We can call this a ‘Nominal Diagnosis’. Here are two examples:

• K, a tense, conscientious, sensitive woman of twenty-six years has seen several doctors over several years with benign abdominal and bowel symptoms. All have agreed she has Irritable Bowel Syndrome (IBS) and prescribed the usual medications, always with little or transient effect. Dr T realises that a wider vocabulary is needed to differently understand and influence her complaints. In his endeavour to do this, he learns a lot about her unhappy childhood home and how this has led to her guarded perfectionism and her painful ambivalence about close relationships. The long-term effects of this widened dialogue and vocabulary were slowly gratifying for both K and her doctor.

Here the conventional noun-diagnosis of IBS was relatively ineffective and – probably – obscuring or obstructing more helpful personal understanding; it
thus proved to be a Nominal Diagnosis only. The idiomorphic understanding she developed with Dr T proved much more helpful.

- Maggie, fifty-five years, has collected a variety of diagnoses from her many years of faltering contact with psychiatrists and psychologists: Generalised Anxiety Disorder, Agoraphobia, Panics, Emotionally Unstable Personality Disorder with Cyclothymia, Recurrent Depressive Disorder, Bipolar Affective Disorder. All are documented in the usual formalised language of designatory healthcare which then rhetorically confine and define Maggie by Nominal Diagnoses. These conferred nouns may superficially appear to offer real therapeutic understanding, leverage and prediction, but actually do not. None have offered Dr V greater personal understanding of Maggie.

Dr V decides to create a larger and different kind of space for Maggie to talk. This dramatically changes not only Dr V’s view and understanding of Maggie, but also Maggie’s behaviour: her symptoms become much quietened.

How does this happen? Dr V wants to know Maggie’s story, not for a Management Plan, but so that he can better understand. Her story has obscurely disturbed her for decades and it will disturb Dr V now.

Twenty-five years ago she was married – happily she thought – with three children. She experienced her husband as kind, attractive and funny, but a bit feckless: he drank a lot. She suddenly has unmistakable evidence of his alcoholicly hazed, repeated sexual contact with their ten-year-old daughter, Amanda. In a volcanic eruption of mixed and primitive feelings, her marriage and family are destroyed. Years later the ruined landscape of her life is still littered with explosives. She tells Dr V of a current torment: Amanda – now a tough, cynical, sexually alluring, drug-abusing, spiky thirty-four year old single mother – has restored affectionate contact with her father and his second wife, and takes her children to see them. Maggie’s feelings towards her daughter are raw, kaleidoscopic and irresolvable: ‘My mind goes crazy with it, doctor … She was only ten: I should have known, should have protected her: but she knew, and she knew that I didn’t know … She was a child, but was – I didn’t know then – a serious sexual rival. Now she is stronger and healthier.
than me, and has more of a family – mine is destroyed. I love her as a mother, but hate her for what she did, what she does, what they have all done: but can I blame her? … Has she triumphed over me? I feel crazy and terrible for having just said all that, doctor, yet it’s such a relief that I can say these things and that you can understand …’

Dr V has a bespoke and ongoing dialogue with Maggie about her tragic story and her responses to it. Her distress often exceeds her capacity for words, yet words and ideas are what they exchange and they are many: guilt, shame, loss, rage, hate, love, contrition, resentment, despair, despondency, alienation, disconnection, blame, humiliation, revenge, sorrow, defeat … All touch on part of Maggie’s poisoned cauldron, but only part, and only transiently. Maggie now talks less of her symptoms and Dr V does not much offer his vocabulary of diagnoses or treatments. As their language has changed, so has the nature of their exchange, and then the pattern of Maggie’s distress. Maggie, so long burdened and defined by her multi-diagnoses, is now freer to suffer with her unique and humanly-understood tragedy. Dr V, too, though distressed by her story, is also touched and, paradoxically, nourished by such candid and courageous contact – the staple of compassion.

Others have asked: what is ‘really’ wrong with Maggie? Is there a word: is it ‘Depression’? What is the ‘correct treatment’?

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3. Never invade Russia!

This was Churchill’s droll, jesting yet ominous response to an enquiry about his most crucial guiding military maxim.

Napoleon and Hitler are the best-known examples of Churchill’s warnings of epic folly: maelstoms of shocking, humbled hubris. Both launched their expeditions fresh from success in easier campaigns and fuelled by specious optimism. They were driven also by rhetoric for the rightness and feasibility for the possession of new territory. Both started with startling triumphalism, then slowed, then succumbed: exhausted by the vastness
and strangeness of a climate, terrain and people they had poorly understood.

There are useful analogies for some of our current enervating endeavours and conundrae in healthcare. First, our expectations have been primed and inflated: Life for millions in the Twentieth Century was positively transformed by applied science. Biomedicine has had spectacular success in countering, even eliminating, many infectious, inflammatory and degenerative physical diseases. In all this industrialisation – mass-production, standardisation, quantification, speed – has been essential. Such successes have led to a long flush of optimism: surely we can gainfully apply similar schematic, industrial-medical type thinking and interventions to all our other sources of distress and pain – our human disease, our polymorphous anguish, our inevitable (yes, still!) decline?

It is here that our invincible march founders, for ailments of our metaphorical heart are proving far harder to locate, define or reverse than those of our anatomical heart. Human motivation, meaning, communication and (un)consciousness yield very meagre territories to objectifying science. Beyond is our vast hinterland, navigable (sometimes) by other kinds of knowledge and influence.

Our reluctance to heed this accounts for many of our most curious and (superficially) indecipherable healthcare follies. In our thrall to measurement we neglect more important unmeasurables. In our urge to treat we do not pause to heal. In our (often unnecessary) compulsion to convergently image the part, we become blind to the divergent – the whole: this person, their story and networks. When we define, we also often confine – ourselves and others – to a tunnelled vision and selective deafness. For language, perception and action are tightly linked. If the language of our culture becomes restricted to the technical, the commercial, the procedural and the defined, then our patients – people, like us! – are seen as merely biomechanical problems to be controlled, managed, traded or disposed of. The abstract becomes hegemonic: the real become abstract.
Hyperbole?

Even in ‘straightforward’ physical care our over-industrialisation is producing shocking calumnies. Consider the following story recently widely reported in the media: 4

- A man is admitted to a London Hospital with a rare but well recognised physical complaint (Diabetes Insipidus) which renders him particularly and hazardously vulnerable to dehydration. He knows this and can usually communicate well. He is seen and assessed by a succession of healthcare workers, some of them specialists. In their complexly successive, jigsawedly interlocking, brief contacts with him they do not heed this increasingly desperate requests for water, which culminate in his calling 999 from the hospital ward. Only after he dies does it become clear that all these algorithmically-managed practitioners had been effectively deaf to his voice and blind to his demeanour. Hospital spokespersons’ public comments are woven with grave contrition and confusion. The former might need construction, the latter does not. The Hospital used to have world-renown for its standards of medical practice, teaching and academia; emblematised also by its historic, stately architecture. Relocated now in an undistinguished, unloved, ugly, airport-like, sprawling conurbation, the containing architecture expresses with unintended accuracy the healthcare culture – a hive of hired healthdroids.

That a highly-funded, well equipped and specialised medical unit can so misunderstand and depersonalise someone with a physical complaint can only bode poorly elsewhere – especially for those who require yet more personal and thoughtful kinds of listening and understanding. This is the case, but often less obvious. With non-physical complaints our failures of care and communication are less dramatic: a slow slide into lonely and dislocated oblivion will gather no headlines. Living silently with a broken-heart attracts no crowds; an untimely death from a heart attack does.

Our current healthcare is in increasing thrall to a Scientistic folly: that generic formulations can be mass produced for all individual distress – that human dis-ease can thus be easily subsumed to impersonally managed
forms of civic engineering. Such is contemporary healthcare’s Invasion of Russia: grandiose but flawed in assumption, then unsustainable, impossible and incurring vast casualties.5

Healthcare may be guided by our science, but science must rarely eclipse our humanity.

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4. If you want good personal healthcare, see a Vet

‘I like peasants – they are not sophisticated enough to reason speciously’

– Montesquieu (1689-1755), Variétés

When Dr F takes his dog to the Vet, Mo, he is simultaneously disarmed, comforted, ashamed and envious: Mo has a guileless and effortless rapport and liking for the animals she is handling. Dr F wants to know more of these unaffected and unbookish skills: he asks to sit in with her.

What Dr F witnesses is humbling and radically refreshing. After asking the owner a few questions, Mo stands back from the animal, scanning it with her eyes, listening carefully to its breathing and other sounds. Then she makes active contact with the animal, the approach being based, Dr F thinks, on some kind of ‘holistic mind-set’ that she senses the animal is now inhabiting. Dr F notices how different her approaches are: with one she gazes at its face with unwavering directness while speaking in a firm and commanding voice; with the next she averts her gaze, softens her posture and lowers her voice to a soft reticence. Sometimes she quickly and directly grasps the nape of the neck with decisive dominance, a wordless control. At others she is slow and light of touch, gently stroking the flank while humming; a trans-species fraternalism. Dr F wonders whether Mo’s accuracy, range and speed of rapport with these different creatures is somehow akin to inducing hypnotic states in humans. He asks Mo:

‘Oh, I don’t know about hypnosis – I’m not that clever. Nor do I know much about humans: they talk too much for me to be able to understand
them!’ She curls a playfully commiserating look at Dr F. ‘My furry friends here can’t say much, but I have to understand them quickly: are they frightened, hungry, confused, in pain, angry, unloved? … Yes, really! … Do they need to feel they still control their territory, or do they need to know I am dominant? All such things I have to get right without much delay, otherwise I cannot get docility enough to do my job … Yes, I’ll get scratched and bitten, too. With larger animals it can be more serious: you can easily become lunch or squash!’.

Dr F leaves Mo that morning with a deeper gratitude than he is easily able to express. With little psychological scholarship, theory or instruction, this open-hearted, open-minded, freshly-instinctive woman is able to resonate with, and thus ‘read’, the mind-set of these (humanly) mute creatures. What natural gifts we (all?) may have!

He thinks of the cumbersome, academically conceived, elaborate-yet-clumsy devices healthcare workers are being instructed to use, to inform all about the experience – the ‘mental state’ – of others. He thinks of the obedient but hopeless Scientism of giving detailed questionnaires to Kenny and Philip. He then thinks of Mo: her almost wordless, seemingly magical, rapid and affectionate rapport with very different animals. He wishes he could be understood like that, and laughs to himself. His laughter diffuses to a smile at the contrasted memories: Mo has inspired him to retrieve some fresh depth and contact in his work. He will reconnect with himself too, before and beyond words.

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5. In difficult encounters, think about sex

The tyranny of the explicit

‘Every person’s feelings have a front-door and a side-door
by which they may be entered.’

Oliver Wendell Holmes Sr, The Autocrat of the Breakfast Table (1858)

Dr Y is thinking about sex. It is not the first time, but now it is different. He is thinking professional thoughts about how our thinking and behaviour around our sexuality could greatly enlighten our healthcare.

More specifically, Dr Y is thinking about a very delicate, complex and evanescent interweaving – of the implicit and the explicit: how these have to be rapidly and accurately discerned, deciphered, jointly understood and then responded to. All of this happens on a second by second basis. And choreographing this medley of meta-communications is essential for any kind of sexual competence – let alone deeper unifying satisfactions. We have to have a (usually) unspoken sense of what the other is desirous of, receptive to, ‘on’ for, and when and how. We must quickly sense error and redirection. Mostly, in better sexual congress, this can happen by dextrous implicit exchanges: the explicit may sometimes then be added potently and sparingly – a mutual aphrodisiac. If the explicit is necessary, the exchange is faltering. If it is necessary for long periods, the relationship is in serious trouble. If the explicit is used by one, without implicit desire by the other, the exchange becomes embarrassed, self-consciously clumsy, possibly abortive. Many such misattunements doom a relationship. Seriously regarded, they can become work for lawyers.

Dr Y is amusing and confusing himself with how weighty and complicated are the responsibilities of this ancient and near universal activity. How do most of us ever (think we) get it right?

These implicit-explicit dances are certainly at the heart of our sexual contacts, but extend throughout our important relationships. They depend on our being able to seamlessly interchange the implicit and explicit, by ‘tuning-in’ to the other. We want (and expect) our partner to understand
what is troubling us, without our having to name it (yet?): soon after, we want them to now be receptive to the beginnings (or resumption) of the explicit. We want, now, to be able to talk. Yes, clearly.

Familiar?

Dr Y extends his thinking to how important such exchanges are in healthcare. He remembers Maggie’s long story and considers how any success he has with her is due to his being mindful of such delicate dances: he had been patiently implicit with her before she trusted him with the explicit. And then, with gratified relief, her healing reverted to the implicit. Maggie had told Dr Y of earlier psychiatric interviews and how they had become too explicit too rapidly. She had retreated to the shelter of the implicit, but had not been understood. The implicit locked.

Dr Y remembers well the kinds of discussions he used to have with colleagues, at the beginning of his career. He recalls many years of interrupted-but-never-finished, free-wheeling explorations of our complex contact with others. The concepts and vocabulary were rich and wide: influence, confluence, identity, boundaries, encryption, territory, projection, surrender, escape … The notions and vocabulary were plastic and uncompleteable, yet each alightment could enrich – differently in different conversations: subtly or evidently, with immediacy or incubation, with implicitness or explicitness.

Dr Y now rarely has such polychromatic and rewarding exchanges. The computer has predicated a new healthcare language for the 21st century: a restricted and restrictive machine-mandated vocabulary. Healthcarers communicate now – almost entirely – in dull narrow administrative, technical words: of conventions, clusters and codes; of quantifiable procedural activity and description; of conduction but not induction – all designating the objectively generic but excluding the humanly variable. Computer compatibility may thus build some bridges to our outer lives, but very few to our inner. What remains has little room for the nascent, the semiotic, the metamorphic, the ambiguous – all the subtle hues that we must mindfully
respect to provide nourishment and meaning for our important relationships. The explicit now burgeons beyond our needs, understanding, tolerance or stamina: the implicit ails and dies.

Its passing takes much of us, too.

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The whole is more than the sum of its (explicit) parts.

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Healthcare is a humanity guided by science.

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Humanity may be commanded by the explicit: its best understanding is often implicit.

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‘The water in the vessel is sparkling; the water in the sea is dark. The small truth has words that are clear; the great truth has great silence.’

– Rabindranath Tagore, Stray Birds (1916)
Notes

1. DWP – a UK nationwide governmental department that administers and manages state pensions, sickness and welfare benefits etc. Assessing disputed levels of distress and disability is a task it currently often subcontracts to other agencies.

2. The doctors, patients and situations throughout this article are real but anonymised. The examples of doctors’ encounters will be relevant now to many other types of healthcare professionals, especially those working for the NHS or large corporations.

3. Detailed questionnaires are now being vaunted and proceduralised throughout most NHS Psychology and Counselling Services. This is explained by authorities as making the services more scientifically efficient. This is contentious, at least. In this author’s view it leads to specious science, dehumanisation, and a healthcare cult of Scientism. The obstructive and destructive effect of these is extensive and subtle. See my articles ‘How to Help Harry’ (Zigmond, 2012) and ‘Sense and Sensibility’ (Zigmond, 2011a).

4. It is hard to gain statistics about the human and economic cost of inflexible, officious practice and uncompassionate – if ‘correct’ – depersonalised care. What correlates does one measure? Who is going to fund this? Although quantitative research may be difficult to set up, vernacular evidence is plentiful and ubiquitous. See my articles ‘Five Executive Follies’ (Zigmond, 2011b) and ‘Love’s Labour’s Lost’ (Zigmond, 2010). Also letters to Clinical Directors of Mental Health Services.

5. Kane Gorny, age twenty-two, died on 25 May 2009 of dehydration as an in-patient at St George’s Hospital, south London. The inquest in July 2012 revealed the facts recorded here. The story is only one of several similar in recent years, eg see also reports of Mid Staffs and West Midlands NHS Trusts. All have been met with convulsions of outraged incomprehension when made public. The fact that they come clearly to public view reflects well on investigative journalism but – of course – seriously damages confidence in NHS care. This is rendered more confusing when such episodes occur in institutions deemed to be ‘performing’ excellently by other, measured criteria. The responses of managerial gravitas, concern and contrition seem real enough. Some sceptics have averred that these conceal some kind of collusion, albeit unconscious. The latter possibility is easier to cite and sense than see. If true, this is cultural: powerful, but difficult to tether or examine, except by inference.

6. The activity itself is much older than many people realise. For example, this author – together with many of his generation – believed they were its initiators in the 1960s. However, since that time there has been increasing evidence from many sources, indicating that it far predates that period – possibly even prior to the birth of this author’s own parents.
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All the above, plus the letters referred to in note 4, are available online at www.marco-learningsystems.com/pages/david-zigmond/david-zigmond.htm

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