MEDICINE AND SOCIETY

What Should Clinicians Do to Engage the Public About Gene Editing?

Tom Alsaigh, MD, Laura Nicholson, MD, PhD, and Eric Topol, MD

Abstract

Genome editing holds tremendous promise for preventing, ameliorating, or even curing disease, but a thorough discussion of its bioethical and social implications is necessary to protect humankind against harm, a central tenet of the original Hippocratic Oath. It is therefore essential that medical students, physicians, and all health care workers have a working understanding of what gene editing entails, the controversy surrounding its use, and its far-reaching clinical and ethical implications.

Gene Editing’s Promise

“Have no fear of perfection—you’ll never reach it.” Arguably one of the closest scientific challenges to Salvador Dali’s famed proclamation came in 2012, when scientists repurposed a bacterial adaptive immune system to make precise edits to genomic DNA with astounding ease and efficiency. The concept of genetic engineering to modify genes has been around since the 1970s, but only relatively recently has its promise materialized due to the discovery of sophisticated gene editing systems. Although multiple forms of gene editing have been studied and refined (eg, zinc-finger nucleases and transcription activator-like effector nucleases), the clustered regularly interspaced short palindromic repeats (CRISPR)/Cas9 gene editing tools have bolstered the promise of correcting genetic miscues due to the relative ease and efficiency with which they can be used. The promise of Cas9, a bacterial-derived DNA-editing enzyme, is its ability to home in on a specific DNA sequence by using a CRISPR RNA guide sequence that is complementary to the target DNA sequence and that binds by Watson-Crick base pairing, thereby allowing Cas9 to cleave the sequence of interest. A CRISPR/Cas9 complex targeting the mutated Huntington gene, for example, could locate the defective DNA sequence and cut it with high accuracy, preventing production of defective Huntingtin protein. This technology has advanced rapidly, and a variety of other genomic modifications are now being introduced by mRNA editing and alternative splicing.

Current Applications

Today, numerous academic and biotechnology groups are focused on the translation of CRISPR/Cas9 technology to correct a variety of genetic diseases. Derivation of induced pluripotent stem cells from patients with the targeted disease is a common model to study the capabilities of CRISPR. By targeting disease-causing mutations, somatic and
germline gene editing could soon be a clinical reality for patients suffering from a variety of diseases, such as β-thalassemia, hemophilia A, cystic fibrosis, Duchenne muscular dystrophy, α1-antitrypsin deficiency, polycythemia vera, HIV-1, and Epstein-Barr Virus.

In addition to preclinical work in gene editing, clinical trials are underway to target aberrantly expressed genes in a variety of disease processes, including malignancy. The University of Pennsylvania is currently enrolling patients for a phase I clinical trial to CRISPR-edit autologous T-cells ex vivo in an effort to target an immunogenic tumor antigen (NY-ESO-1) in relapsed refractory multiple myeloma, melanoma, synovial sarcoma, and myxoid/round cell liposarcoma. In late 2018, the pharmaceutical company Editas Medicine received approval from the US Food and Drug Administration for a phase I/II in vivo trial to correct a point mutation in the CEP290 gene for Leber’s congenital amaurosis type 10, the most common form of inherited childhood blindness.

Finally, due to the shortage of organs available for transplantation, xenotransplantation—the transfer of living cells, tissues, or organs from one species to another—is being reconsidered due to advances in CRISPR editing. Specifically, scientists have used CRISPR/Cas9 to inactivate porcine endogenous retroviruses, thereby preventing cross-species transmission and mitigating harmful pig-to-human immune incompatibilities. These examples demonstrate the extraordinary promise of human gene editing, which raises exciting possibilities for treating a plethora of diseases but also introduces a variety of ethical and societal challenges.

Somatic vs Germline Gene Editing
An important distinction that will help guide our discussion of ethical implications of human gene editing is the difference between somatic and germline gene editing. Somatic gene editing involves making nonheritable genetic modifications to a person’s genome to treat the disease being targeted. By contrast, in germline gene editing, egg or sperm DNA is altered, and these modifications affect all subsequent cell types following fertilization and are transmitted to future generations, potentially altering the gene pool. The heritability of germline gene alterations is one of the main reasons why the World Health Organization (WHO) is developing an advisory panel to oversee and provide guidance on human gene editing, and it highlights the importance of reaching international consensus on appropriate uses of germline editing technologies.

Managing Risk and Hope
In 2015, just 3 years after CRISPR-Cas9 was used to cleave DNA in vitro, the Economist brought to life in its cover issue what many feared would become a reality with the advent of germline gene editing: eugenics via the creation of “designer babies” with enhanced features and characteristics. In the United States, 72% of respondents polled in 2018 said that gene editing is an appropriate use of technology to treat a serious
disease or condition. But only 19% answered in the affirmative when asked whether this technology should be used to make babies more intelligent, underscoring the idea that these efforts should focus on treating serious illnesses.

By late 2018, gene editing suddenly faced blistering criticism when He Jiankui of China announced that, for the first time, human germline gene editing had been used to confer 

HIV resistance by modifying the CCR5 gene in embryos that were then implanted, 

producing twins. The choice to use germline gene editing for the purpose of preventing 

HIV transmission was highly controversial, as some have argued that the focus of 

human gene editing should be to serve an unmet clinical need, and the US Department of 

Health and Human Services notes that the risk of the virus being transmitted from 

mother to baby is 1% or less when a pregnant woman is treated appropriately. 

Furthermore, experiments done to render the twins immune to HIV could cause them 

serious harm. Some studies suggest that deletion of the CCR5 gene can potentially 

increase susceptibility to West Nile virus and tickborne encephalitis and have 

additional deleterious effects on immune responses. Perhaps even more importantly, 

the demonstration that this technology was viable in human applications without the 

risks being first fully considered opened up Pandora’s Box.

In March 2019, several leading gene editing experts called for a global moratorium on 

germline genome editing in humans in order to give the international community time 

to establish a more detailed framework by which to guide its future use. Part of the need 

for such a moratorium is the persistence of many uncertainties about the consequences 

of gene editing, such as off-target effects, or unintended cleavages of DNA sequences. 

There is also the chance that only some copies of targeted genes are modified, causing 

mosaicism.

So how should we, as medical professionals, address these concerns, and which values 

should guide clinical and research practice? In order to mitigate the potential adverse 

effects of gene editing, we need well-designed preclinical studies that support uses of 

gene editing for patients’ unmet clinical needs. Of utmost importance are the scientific 

rigor with which these studies are evaluated and the publication of both positive and 

negative findings. Every proposal for gene editing in human embryos that would not be 

brought to term should be subject to rigorous international oversight, even during study 

design, to ensure proper informed consent and high technical standards that motivate 

scientific rigor and integrity.

Because gene editing is enticing and has now actually been done in humans, 

organizations such as the National Academy of Sciences, Engineering and Medicine 

(NASEM) have set forth principles to guide somatic and germline gene editing in clinical 

practice and human subjects research (see Table).
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<th>Principles</th>
<th>Description</th>
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<tr>
<td>Promote well-being</td>
<td>“Providing benefit and preventing harm to those affected.”</td>
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<tr>
<td>Transparency</td>
<td>“Openness and sharing of information in ways that are accessible and understandable” to patients, their families, and other stakeholders.</td>
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<td>Due care</td>
<td>Proceeding with research “only when supported by sufficient and robust evidence.”</td>
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<tr>
<td>Responsible science</td>
<td>Adhering “to the highest standards of research ... in accordance with international and professional norms.”</td>
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<tr>
<td>Respect for persons</td>
<td>Recognizing “the personal dignity of all individuals ... and respect for individual decisions.”</td>
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<td>Fairness</td>
<td>Treating all cases alike, with an equitable distribution of risks and benefits.</td>
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<tr>
<td>Transnational cooperation</td>
<td>Committing “to collaborative approaches to research and governance while respecting different cultural contexts.”</td>
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*Adapted from National Academies of Sciences, Engineering and Medicine.*

**Individual and Collective Impact**

Making decisions and contributing ideas during decision making processes are important expressions of autonomy, a value that both practitioners and patients hold sacred. One central problem with human germline gene editing is that autonomy is taken away from an individual, even before birth. In a poignant piece published in *Nature*, a young girl with albinism and blindness, full of determination to play soccer, was asked if she wished her parents had corrected the genes that contributed to her genetic condition before she was born. Without a second thought she answered no. In a thought-provoking twist, their having done so might have changed her character or ambition or resulted in her being less motivated to overcome challenges. Were this question asked of someone else, that person might answer differently than she did, but it is presumptuous to think that every person would want genetic “defects” to be edited prior to birth. The perspective of persons with disabilities gives credence to the idea that germline gene editing might best serve society if used to prevent serious illnesses or fatal germline inheritance. A discussion about what separates serious from nonserious illness is perhaps an important one to have.

Because an unborn person is unable to participate in these kinds of conversations, there should be broad societal consensus about what constitutes acceptable use of germline gene editing. Furthermore, because members of different cultures hold different values, international governance of human gene editing is complex. While the aforementioned
moratorium on human germline gene editing outlines key issues that should be addressed before resuming human germline gene editing experimentation, it also expresses respect for sovereign nations’ opting to resume experimentation if certain criteria are met, such as engaging the public, offering justification for national implementation plans, and attaining societal consensus before proceeding.37

Engagement With Patients and the Public
The NASEM and other scholars have identified strategies that clinicians can use to engage patients and other stakeholders in discussions of human gene editing.38,41 Communication about gene editing and its nuances should reach a broad cross-section of society, including advocacy groups, religious communities, and well and poorly educated segments of the population. Clinicians and health care workers should be able to explain the basics of gene editing and its potential uses in health care. Clinicians’ consultation with advocacy groups could also help spread information, as these groups can continue broader discussions of pertinent topics among their stakeholders. Finally, exchanges of decision-relevant information through dialogue can increase the spread of helpful information and awareness among the public. By using our platform as clinicians who care for a broad cross-section of society on a daily basis, we can help guide, inform, and grow public conversations about gene editing.

Conclusion
Despite the sometimes-negative media attention to human gene editing, scientific curiosity and discovery should not be stifled. Without bold ideas accompanied by good intentions, we would not have great scientific discoveries, such as vaccination or organ transplantation. However, bold ideas must also be accompanied by rigorous regulation to guarantee transparency, the ethical conduct and beneficial intention of gene editing studies, and protection of vulnerable patients and communities. Ultimately, as a society, we must try to carefully distinguish what is medically necessary from what is medically or socially desirable. Clinicians can have central roles in shaping these conversations.

References


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