Towards the elements of successful insect RNAi

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Abstract
RNA interference (RNAi), the sequence-specific suppression of gene expression, offers great opportunities for insect science, especially to analyze gene function, manage pest populations, and reduce disease pathogens. The accumulating body of literature on insect RNAi has revealed that the efficiency of RNAi varies between different species, the mode of RNAi delivery, and the genes being targeted. There is also variation in the duration of transcript suppression. At present, we have a limited capacity to predict the ideal experimental strategy for RNAi of a particular gene/insect because of our incomplete understanding of whether and how the RNAi signal is amplified and spread among insect cells. Consequently, development of the optimal RNAi protocols is a highly empirical process. This limitation can be relieved by systematic analysis of the molecular physiological basis of RNAi mechanisms in insects. An enhanced conceptual understanding of RNAi function in insects will facilitate the application of RNAi for dissection of gene function, and to fast-track the application of RNAi to both control pests and develop effective methods to protect beneficial insects and non-insect arthropods, particularly the honey bee (Apis mellifera) and cultured Pacific white shrimp (Litopenaeus vannamei) from viral and parasitic diseases.

Keywords
Antiviral therapy, dsRNA, Insect pest control, RNA interference, siRNA, Systemic RNAi

Disciplines
Biology | Entomology | Genetics

Comments
This article is from Journal of Insect Physiology 59 (2013): 1212–121, doi:10.1016/j.jinsphys.2013.08.014.

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RNA interference (RNAi), the sequence-specific suppression of gene expression, offers great opportunities for insect science, especially to analyze gene function, manage pest populations, and reduce disease pathogens. The accumulating body of literature on insect RNAi has revealed that the efficiency of RNAi varies between different species, the mode of RNAi delivery, and the genes being targeted. There is also variation in the duration of transcript suppression. At present, we have a limited capacity to predict the ideal experimental strategy for RNAi of a particular gene/insect because of our incomplete understanding of whether and how the RNAi signal is amplified and spread among insect cells. Consequently, development of the optimal RNAi protocols is a highly empirical process. This limitation can be relieved by systematic analysis of the molecular physiological basis of RNAi mechanisms in insects. An enhanced conceptual understanding of RNAi function in insects will facilitate the application of RNAi for dissection of gene function, and to fast-track the application of RNAi to both control pests and develop effective methods to protect beneficial insects and non-insect arthropods, particularly the honey bee (Apis mellifera) and cultured Pacific white shrimp (Litopenaeus vannamei) from viral and parasitic diseases.
1. Introduction

RNA interference (RNAi) has transformed insect science research because it enables the researcher to suppress a gene of interest and thereby link a phenotype to gene function. For basic research purposes, RNAi offers a route to functional genetics in all insects, including those for which transgene resources do not exist (Belles, 2010). RNAi also has enormous potential for applied entomology (Price and Gatehouse, 2008; Xue et al., 2012). For example, RNAi can be used for insect pest control by suppressing essential genes leading to reduced fitness and/or mortality. Furthermore, by priming the antiviral RNAi response with innocuous viral sequences, beneficial insect species, such as honey bee (Apis mellifera) and silkworm (Bombyx mori), can be protected from highly pathogenic viral infections. However, the reality is not yet matching the envisioned potential of RNAi. Practitioners are increasingly aware that RNAi in insects can be capricious; efficacy varies across insect taxa, among genes, with mode of delivery, and even between different laboratories (Terenius et al., 2011). All too often, the application of RNAi technology is an empirical exercise: “try it, for it might work”.

The goal of this article is to promote the use of practical principles to design and interpret insect RNAi studies. We know that no single protocol can be applied for every gene in every insect. Therefore, the specific purpose of this article is to provide a roadmap for the application of RNAi for experimental analysis of gene function, management of pests and protection of beneficial arthropods. The article is divided into three sections. First, current knowledge of the mechanisms and function of RNAi in insects is reviewed, highlighting the known variation among insect taxa. This information offers a guide to the most appropriate strategy for different insect systems, and provides the springboard for much-needed future innovation in RNAi technology. Second, the design of RNAi studies is addressed, using both empirical data and conceptual understanding to identify successful experimental designs, effective methods for RNAi delivery, and informative indices of RNAi efficacy. Importantly, there is no single protocol for the perfect RNAi experiment, partly because the efficacy of RNAi strategies varies among insect groups. In the third section, we turn to the application of RNAi for the management of pest and beneficial insects, and discuss the unique opportunities and challenges associated with each of these applications.

2. Mechanisms of RNAi

RNAi refers to the suppression of gene expression by small non-coding RNA molecules, predominantly by the cleavage of a target mRNA in a sequence-specific manner (Fire et al., 1998), and the general steps involved in this process are shown in Fig. 1. Upon cell entry and recognition, double stranded RNA (dsRNA) is cleaved by the RNase III Dicer into 20–25 bp fragments with a two base overhang at the 3’ end. These fragments are incorporated into the multi-protein RNA-induced silencing complex (RISC), where one strand (the “passenger” strand) is eliminated and the other “guide” strand is retained. The catalytic component of RISC is the RNase H-like domain of an Argonaute protein, which cleaves single-stranded RNA molecules having sequence complementary to the guide RNA. Most eukaryotes, including animals and plants, have Dicer and Argonaute proteins, and possess the RNAi machinery (Shabalina and Koonin, 2008).

The 20–25 bp RNAs generated by Dicer comprise two groups (Ghildiyal and Zamore, 2009; Matranga and Zamore, 2007; Asgari, 2013): microRNAs (miRNAs), which are processed from endogenous gene transcripts and function in the regulation of gene expression, and small interfering RNAs (siRNAs), which are derived from dsRNA molecules and provide defense against viruses and transposable elements. The experimental use of RNAi exploits the siRNA pathway, specifically the capacity of cells to degrade a single-stranded RNA (ssRNA) (including mRNAs) with sequence identity to the administered dsRNA molecules.

Three processes determine what can be achieved by RNAi: cellular uptake of the RNAi molecule (usually dsRNA), the production of secondary dsRNA molecules in the cell, and the transfer of these molecules to other cells (Fig. 1). Where the RNAi–mediated silencing is transmitted widely throughout the treated organism, RNAi is described as systemic (Whangbo and Hunter, 2008; Huvenne and Smagghe, 2010). In principle, the success of an RNAi experiment could be predicted from the level of these activities in the insect of interest, and strategies that increase these activities...
might enhance RNAi-mediated knockdown of the target gene expression. The difficulty is that we have little or no understanding of whether or how dsRNAs are amplified within insect cells or disseminated among insect cells.

In C. elegans, systemic spread of RNAi is optimal for dsRNA molecules $\geq 50$ bp long, and is independent of Argonaute function (Feinberg and Hunter, 2003; Tabara et al., 1999), suggesting that the molecules moving between cells are dsRNAs, and not amplified siRNA. Uptake of dsRNA by somatic cells requires the protein SID-1 (systemic interference defective-1), which is inferred to function as a dsRNA channel (Winston et al., 2002). Other C. elegans proteins, SID-2 and SID-5, have been implicated in dsRNA uptake by gut cells and dsRNA export from cells, respectively (Hinas et al., 2012; Winston et al., 2007). Putative insect orthologs of the C. elegans sid genes have been described (e.g. Dong and Friedrich, 2005; Xu and Han, 2008), but these reports deserve careful evaluation. The sid-1-like genes in various insects have a greater sequence identity with the C. elegans gene chup-1, also known as tag-130, than to sid-1. CHUP-1 is a cholesterol transporter and has no known involvement in RNAi (Valdes et al., 2012). As an illustration, a gene in Locusta migratoria initially identified as sid-1-like is not required for systemic RNAi (Luo et al., 2012), and is the ortholog of C. elegans CHUP-1 (as determined by top reciprocal BLASTp hit), and not SID-1.

Plants and the nematode Caenorhabditis elegans possess an RNA-dependent RNA polymerase (RdRP) (Pak and Fire, 2007; Xie et al., 2001) that facilitates the within cell amplification of silencing. The fragments of the target ssRNA released from the RISC act as a template for RdRP-dependent dsRNA synthesis, yielding more substrate for RISC-mediated degradation of the target ssRNA. Efficient amplification of RNAi by RdRP can drive the abundance of the target ssRNA molecule to undetectable levels, and RdRP is essential for RNAi in C. elegans (Sijen et al., 2001). RdRP has been identified in a few animal species beyond Caenorhabditis nematodes, including the cephalochordate Branchiostoma floridana (Vienne et al., 2003), but no verified RdRP homolog is evident in any insect genome sequenced to date (Tomoyasu et al., 2008). It is unclear whether or how the RNAi triggered by the acquisition of dsRNA molecules is sustained in insect cells (Fig. 1).

Analysis of gene orthology between C. elegans and insects has been a productive approach to identify the core RNAi machinery in insects, but far less informative for understanding the molecular basis of intracellular amplification and systemic spread of RNAi. The alternative discovery-based strategy of a genetic screen, using rescue from RNAi-lethality, has great potential. For example, Ulvila et al. (2006) demonstrated that uptake of dsRNA by Drosophila S2 cells is strongly endocytosis-dependent, and mediated principally by the scavenger receptors Eater and SR-CI. We should, however, be cautious in extrapolating from these data to organismal RNAi because S2 cells, which are hemocyte-like, display high rates of endocytosis as compared to the majority of cell types in the intact insect.

The physiological role of insect RNAi could be informative in predicting RNAi efficacy in different insect taxa and cell types and, by extension, the development of sustainable strategies for RNAi applications in field conditions. There is a strong consensus that RNAi contributes to insect immunity against viruses with a dsRNA genome or dsRNA replicative intermediates in the cytoplasm of infected cells (Blair, 2011; Schnettler et al., 2012). We could, therefore, expect greatest success with RNAi in insect species and cell types that utilize RNAi as a primary anti-viral immune response. Some insects or cell types may have low responsiveness to exogenously-applied dsRNA because they utilize alternative anti-viral defenses (e.g. apoptosis of infected cells, symbiont-mediated protection) (Merkling and van Rij, 2013). It should also be noted that certain insect viruses suppress RNAi. For example, flock house virus (FHV) codes for a protein, known as the B2 protein, that binds to dsRNA, including the FHV replication intermediate, preventing cleavage by insect Dicer and incorporation into RISC (Chao et al., 2005). An insect infected with an asymptomatic, persistent virus that codes for an RNAi suppressor would display limited responsiveness to experimental RNAi (Berry et al., 2009).

3. Designing a RNAi experiment

As described above, RNAi application and efficacy remains variable between genes, organisms and life stages, despite the tremendous utility that RNAi presents for improving our understanding of fundamental biological questions and for pest control. In addition, in insect species where RNAi is predominantly environmental with little evidence for systemic propagation, interference can vary widely between tissues due to differences in the efficacy of dsRNA uptake. Extreme examples are D. melanogaster and Manduca sexta where transcript knockdown by injection of dsRNA has only been achieved in hemocytes, which are capable of endocytosis (Miller et al., 2008; Terenius et al., 2011). In mosquitoes, most tissues can be reached by the injection of dsRNA, however the success of knockdown in the central nervous system varies highly between genes and may be dose-dependent (Lycett et al., 2006; Biessmann et al., 2010). Tissue differences in RNAi efficacy may be overcome by the design of new delivery methods, including transgenesis or viral transduction, which eliminate the requirement for cellular uptake of the RNAi trigger. Development of such technologies is lacking for the majority of species (Fig. 2 and Section 3.2 below).

The aforementioned biological variables, including presence/absence of the core RNAi machinery, cellular uptake and propagation of signal (Rognant et al., 2003; Miller et al., 2008), and dsRNA degrading enzymes (Arimatsu et al., 2007), as well as other differences in genetic backgrounds (Kitzmann et al., 2013), greatly affect
the success of RNAi experiments in different species. Often, these
challenges can be mitigated by experimental factors including
the design of the RNAi molecule, the mode of delivery and the dose
of the dsRNA molecule.

3.1. The RNAi molecule

The success of an RNAi experiment hinges on the production of a
specific RNAi molecule (in the form of dsRNA, siRNA, or a hairpin
RNA) for a target gene of interest (GOI). Experiments should in-
clude an RNAi molecule against a heterologous sequence absent
from the target insect’s genome (typically green fluorescent pro-
tein (GFP) or LacZ), to control for both the administration of the
experimental dsRNA and the physiological impact of triggering
the RNAi cascade. In some cases, a positive control can be incorpo-
rated into the experimental design. For example, in Tribolium, the
use of RNAi against vermillion (white) or Lac-2 provides rapid phe-
notypic evidence of RNAi success manifest in white compound
eyes or white pupae, respectively (Arakane et al., 2005, 2011).

A crucial consideration is the choice of sequence for dsRNA
preparation, especially its length and sequence identity to the tar-
target transcript of the insect. Huvenne and Smagghe (2010) provide
a comprehensive survey of the length range of dsRNAs used in early
studies: from 134 to 1842 bp, with most studies using 300-520 bp.
Comparisons among gene regions (e.g., 5’ end) to which RNAi mol-
ecules are designed have yielded variable results. For example,
RNAi against hunchback (hb) in Acryrthosiphon pisum resulted in
similar mortality whether the RNAi trigger was designed against
the 5’ or 3’ end of the gene (Mao and Zeng 2012), but the 3’ portion
of the inhibitor of apoptosis gene in Aedes aegypti yielded a greater
effect on mosquito mortality than dsRNA targeting the 5’ or central
region of the gene (Pridgeon et al., 2008), and the most effective
antiviral RNAi molecule against infectious myonecrosis virus
(which infects the Pacific white shrimp, Litopenaeus vannamei)
was at the extreme 5’ end of the genome (Loy et al., 2012). These
varied results illustrate the importance of screening multiple RNAi
sequences for a gene of interest.

Generally speaking, greater success with insect RNAi has been
obtained with dsRNA molecules ≥50–200 bp in length (Huvenne
and Smagghe, 2010), although the minimal length required to ob-
tain maximal biological activity varies among insect species
(Bolognesi et al., 2012). Suppression of gene expression has been achieved
with siRNAs (either synthesized directly or obtained by “dicing” the dsRNA in vitro before administration to the insect),
for example in the lepidopteran Helicoverpa armigera (Kumar
et al., 2012), aphid A. pisum (Mutti et al., 2006) and tsetse (Attardo
et al., 2012). It may, sometimes, be necessary to design the RNAi
molecule of shorter length than ideal to obtain specificity, es-
pecially where one member in a gene family that has high sequence
similarity is being targeted. Regardless of the desired size of the
RNAi molecule, the design process can be aided by software that
is informed by genome sequence and RNA folding kinetics to opti-
mize effectiveness; for example, E-RNAi currently offers dsRNA
and siRNA design suggestions for A. mellifera, Tribolium castaneum,
A. pisum, Anopheles gambiae and Ae. aegypti (Horn and Boutros,
2010).

A further issue to be considered in the design of RNAi molecules
is the exquisite specificity of RNAi. In the context of field applica-
tions of RNAi, this property facilitates designing insect-letal se-
quences that are highly species-specific. For example, feeding four
species of Drosophila with species-specific vATPase dsRNA resulted
in reduced vATPase mRNA and significant mortality in conspecific,
but not heterospecific flies (Whyard et al., 2009). In basic research
pursuits, this property affords researchers the capacity to silence
alleles (using short dsRNA) of the same gene specifically, e.g. TEP1
alleles in An. gambiae (Blandin et al., 2009). Conversely, two alleles
of a heterozygous individual, as well as genetically-distinct mem-
bers within an insect population, whether in the laboratory or field,
may differ in their susceptibility to RNAi. This concern is amply jus-
tified by studies on the effect of mismatches between the dsRNA
and its intended target (i.e. mRNA) using synthetic siRNAs admin-
tered to mammalian cells in culture. Most single mismatches
impair the RNAi effect (Birmingham et al., 2006; Jackson et al.,
2003; Joseph and Osman, 2012a,b; Wu et al., 2011); some mis-
matches, however, alter the cellular response from one of transcript
loss (siRNA) to translational repression, (Hu et al., 2010; Tomari
et al., 2007). The advantages of using longer >200 bp dsRNA for
RNAi strategies in pest management is the production of many
siRNAs against the targeted mRNA transcript; potentially maximiz-
ing the RNAi response. Further studies will be necessary to clarify
the extent to which the responses to mismatches in dsRNA and
target mRNA in whole insects differ from the siRNA studies which
used a single construct conducted on cultured mammalian cells.

3.2. RNAi delivery

Efficacy of an RNAi experiment can be influenced strongly by
the mode of delivery of the RNAi trigger (Fig. 2 and references
within). The most widely used routes for administering RNAi to
sects are injection into the hemolymph and feeding. Microinjection
was used in the first successful application of RNAi in an insect, to
obtain knockdown of frizzled in Drosophila melanogaster (Kenner-
dell and Carthew, 1998). This method was quickly transferred to
T. castaneum (Brown et al., 1999) and subsequently applied to adult
insects in An. gambiae (Blandin et al., 2002). Microinjection has
been applied to all life stages in hemi- and holometabolous insects
in a rapidly growing number of orders; indeed routine protocols
are now in place for injection for various taxa, including Tribolium,
B. mori, several genera of Diptera, the honey bee, cockroaches and
orthopterans [for a list of references, see Belles (2010)].

An important barrier to the use of microinjection in some in-
ssects is non-specific damage caused by mechanical damage, which
is most often pronounced when targeting embryos. Experimental
variables that influence survivorship include methods of immobil-
ization (cold, CO2, adherence to a substrate), injection volume, site
of injection, and diluents. Although water or physiologic saline
work well for most species, the diluent may require adjustment
to the particular osmotic pressure of the hemolymph.

Oral delivery is a less-invasive and potentially a high-through-
put method for RNAi delivery. It has particular value for insects
that are intolerant of injection (Fig. 2) and for field applications
for RNAi-mediated pest control (see Section 4). Protocols for
administration of dsRNA synthesized in vitro and incorporated
into the diet are now available for honey bees, aphids, whiteflies
and psyllids (Aronstein et al., 2006; Wuriyangkanhan et al., 2011;
Ghanim et al., 2007; Whyard et al., 2009). RNAi delivery to phytophagous
insects can also be achieved by engineering plants to express
dsRNAs in plant systems for which transgene introduction technol-
ogies are available (Fig. 2). Two complementary methods are in
use: stable transformation by hairpin dsRNAs that target insect
genes (Baum et al., 2007) and transient virus-induced gene silenc-
ing (VIGS), in which engineered viral vectors carrying the gene se-
cuence of interest are transformed into Agrobacterium tumefaciens
and infiltrated into the plant tissue (Burch-Smith et al., 2004). Both
approaches have been exploited, to achieve transcript suppression
in Coleoptera (Baum et al., 2007), Lepidoptera (Baum et al., 2007;
Kumar et al., 2012) and Hemiptera (Pitino et al. 2011; Zha et al.,
2011). In some species, notably dipterans, oral delivery of RNAi
triggers has yielded less consistent results than microinjection
(Zhang et al., 2010). Further, in Lepidoptera, feeding as a mode of
delivery necessitates the provision of high doses of RNAi trigger
(Terenius et al., 2011). This can be attributed to a variety of factors.
The efficacy of RNAi of midgut transcripts may be reduced due to low or inconsistent doses taken up by individual insects, frequency and size of feeding, plus GI tract morphology and physiology will affect the actual dose of RNAi that reaches the midgut epithelium. In addition, there is evidence for production of mRNAs that encode putative secretory dsRNA-degrading enzymes in insects, notably *B. mori*, that can interfere with the RNAi response ([Arimatsu et al., 2007; Liu et al., 2012]). Establishing protocols for consistent RNAi induction by feeding in different species may, therefore, prove challenging. In addition, oral delivery of RNAi molecules in species where systemic RNAi cannot be achieved limits its application to genes expressed in gut cells (Fig. 2).

A minority of studies have exploited alternative routes for dsRNA delivery, including electroporation, soaking or ectopic application, incorporation into nanoparticles, expression in bacteria, topical application, injection into woody plants, direct absorption of dsRNA in water solution into plant cuttings, or root seedlings and trees and solubilization using transfection agents, such as Lipofectamine™ ([Wang et al., 2011; Karim et al., 2010; Zhang et al., 2010; Prigeon et al., 2008; Lopez-Martinez et al., 2012; Hunter et al., 2012]). For additional consideration of this topic, the reader is referred to Yu et al. (2013), which provides a comprehensive review of the history and current practice.

### 3.3. RNAi dosage

The requisite dose of RNAi molecules varies with insect species, life stage, the target gene transcript abundance and its spatial and temporal expression profiles, and according to the delivery method of choice. The viscosity of high dsRNA concentrations limits the injectable concentrations to 6 μg μl⁻¹ (K. Michel, unpub data), and the cost of synthesizing large amounts of dsRNA presents a challenge for high concentrations in artificial diets. Species- and tissue-specific biological factors, including the degradation of dsRNA, and weak activity of the RNAi machinery, can influence the efficacy of RNAi, often requiring relatively high dosage of RNAi molecules. There is now persuasive evidence for dsRNase activity in various extracellular fluids of insects, including the digestive juices of *B. mori* ([Arimatsu et al., 2007]), the salivary of the hemipteran *Linus lineolaris* ([Allen and Walker, 2012]) and the hemolymph of *M. sexta* ([Garbutt et al., 2013]). Although, to our knowledge, this has not been reported in insects, the difficulties in achieving RNAi of genes expressed in neurons of the nematode *C. elegans* has been attributed to the high expression of a nuclease (eri-1, enhanced RNAi-1) in these cells ([Kennedy et al., 2004]).

The mode of uptake, ability to spread RNAi molecules and ability to process the RNAi molecules are other important considerations that no doubt strongly influence the requisite dose required to induce a RNAi response. In *D. melanogaster* larvae, cell autonomous RNAi can be induced readily by the expression of short hairpin RNAs from a transgene; however, injected dsRNAs fail to trigger RNAi in most tissues with the exception of hemocytes ([Miller et al., 2008]). A higher dose is usually required when the RNA molecule is delivered orally as compared to injection. Multiple introductions of dsRNA can enhance the efficacy of RNAi in the salivary glands of *Rhodnius prolixus* ([Araujo et al., 2006]), and although the basis for this effect is not fully understood, one attractive hypothesis is that elements of the RNAi machinery may be expressed at low levels in some tissues ([Chintapalli et al., 2007; Rinkevich and Scott, 2013]), but can be induced in response to the RNAi molecule ([Garbutt and Reynolds, 2012; Liu et al., 2013]).

Peritonym, *Ae. aegypti*, mounts an antiviral RNAi response to Sindbis virus infection, but transcript levels for *Dicer* and *Argonaute* do not change appreciably; only Tudor staphylococcal nuclease, an element of the RISC, shows moderate increase in transcript abundance during an active RNAi response ([Campbell et al., 2008]). Further research is required to establish the incidence and significance of inducibility in RNAi function.

### 3.4. Choice of gene: transcript abundance and protein stability

In principle, the ideal gene target for RNAi produces an mRNA pool with high turnover that codes for a protein with a short half-life. The use of RNAi for phenotypic analysis of gene function in any life stage could be more difficult if the protein product of the target gene has a long half-life. For example, nicotinic acetylcholine receptors (nAChRs) can be stable for >2 weeks ([Lomazzo et al., 2011]) and this protein stability may explain the weak phenotypic response associated with RNAi-mediated knockdown of *Dx6* (nicotinic acetylcholine receptor subunit) expression in both *D. melanogaster* and *T. castaneum* ([Rinkevich and Scott, 2013]). However, for the great majority of genes, mRNA turnover and protein half-life are not known. This gap in our knowledge presents a major challenge for RNAi experiments.

### 3.5. Evaluation of RNAi experiments

The desired result of an RNAi experiment varies with the purpose of the study. High insect mortality is a successful outcome for investigations designed to identify novel RNAi-based strategies to control an insect pest, but a hindrance to many experimental investigations of gene function. For many analyses of gene function, physiological indices of predicted function should be central to the analysis. For example, if a gene under study has a predicted role in protein digestion, osmoregulation or olfaction, then analyses of gut protease activity, hemolymph osmotic pressure and electroantennogram data, respectively, may be useful physiological indices. For some experiments, it may be necessary to reduce the RNAi dose to obtain a reliable physiological signal of gene function obtained by an intermediate expression knockdown, because strong knockdown could result in secondary, deleterious effects on insect fitness that obscure the primary lesion. It is, therefore, important to define the appropriate physiological and fitness assays as an integral part of the experimental design.

The successful reduction of transcript levels as a result of RNAi is most commonly measured by RT-qPCR and expressed as a percent reduction of the relevant transcript in the treatment group versus the negative control group (in which animals were subjected to an RNAi molecule for a heterologous gene). Although this methodology is widely accepted, the choice of reference or housekeeping genes for calculating relative transcript levels is challenging. Even if reference genes for RT-qPCR have been described and validated on the species level, the expression of a reference gene may vary with the physiology and the tissue being targeted ([e.g. Ponton et al. (2011) *Drosophila*, Scharlaken et al. (2008) *honey bees*, Majerowicz et al. (2011) *Rhodnius*]).

Ultimately the phenotypic result of an RNAi experiment hinges on the reduction of protein levels for the gene of interest, and it is highly desirable to determine relative protein concentration. The effect of RNAi on the protein may not be well-correlated to the level of transcript suppression. For example, following dsRNA injection targeting *An. gambiae* SRPN2, SRPN2 protein is not detectable by western blot in the hemolymph, but transcript levels remain at 40–60% compared to controls ([Michel et al., 2005]). Finally, it is possible that RNAi could lead to suppression of transcript (and protein), but not yield a phenotype, particularly where redundancy is built into a specific biological function. For example, deletion of one of the most abundant nAChRs in the insect nervous system results in flies that are “normal” ([Perry et al., 2007]). Whether redundancy will present a limitation for a significant number of other genes remains to be established.
4. Application of RNAi for the management of insect populations

The potential of RNAi for the management of pest insects and protection of domesticated beneficial insects, especially the honey bee, is widely recognized (Xue et al., 2012). In principle, the sequence used in RNAi can be tailored to any taxonomic scale, from a single genotype to a family or even order of insects; and the identity of the target sequence can be manipulated at will, enabling the practitioner to respond rapidly to novel pest taxa or to diminishing efficacy (due to the evolution of resistance, for example) of one target sequence or combination of sequences. In other words, RNAi offers exquisite specificity and flexibility that cannot be matched by traditional chemical insecticides, biological control by natural enemies, or plants bearing protein-coding transgenics.

4.1. RNAi and the control of insect pests

Proof of principle for the application of RNAi in insect crop pest control comes from early studies conducted on the western corn rootworm, Diabrotica virgifera virgifera (WCRW) (Baum et al., 2007), and cotton bollworm Helicoverpa armigera (CBW) (Mao et al., 2007). Baum et al. (2007) fed larval WCRW on 290 dsRNAs, from which they identified 14 genes that reduced larval performance, and one of these, vacuolar ATPase subunit A (V-ATPase), was carried forward for detailed analysis. Low concentrations of orally-delivered dsRNA against V-ATPase in artificial diet suppressed the corresponding WCRW mRNA. Importantly, larvae reared on transformed corn plants that express V-ATPase dsRNA also displayed reduced expression of the V-ATPase gene and caused much reduced plant root damage (Baum et al., 2007). In the study of Mao et al. (2007) on CBW, the target gene was a cytochrome P450, CYP6AE14, which is expressed in the larval midgut and detoxifies gossypol, a secondary metabolite common to cotton plants. When CBW was exposed to either Arabidopsis thaliana or Nicotiana tabacum expressing CYP6AE14 dsRNA, levels of this transcript in the insect midgut decreased, larval growth was retarded, and both effects were more dramatic in the presence of gossypol (Mao et al., 2007). Transgenic cotton plants expressing CYP6AE14 dsRNA also support drastically retarded growth of the CBW larvae, and suffered less CBW damage than control plants (Mao et al., 2011). The research on both WCRW and CBW has been extended to additional genes. The Snf7 gene, which is involved in trafficking of membrane receptors, has been reported to be effective against both D. v. virgifera and D. v. Howardii larvae (Bolognesi et al., 2012; Ramaseshadri et al., 2013); and cotton plants engineered to express cysteine proteases attenuated the peritrophic matrix of CBW, resulting in increased uptake of the dsRNA (Mao et al., 2013). Importantly, cotton plants expressing both the dsCYP6AE14 and cysteine protease were more protected from bollworm than either of the single-transgene lines (Mao et al., 2013).

The studies of Baum et al. (2007) and Mao et al. (2007) illustrate two key issues for successful RNAi of insect crop pests: choice of the target sequence(s) for RNAi; and mode of delivery. The target gene must be an essential insect gene that is consistently expressed through the relevant life-stages and yields reliable RNAi-induced depression of insect performance. As the technology moves from proof of principle to application, very careful consideration of the design of the target sequence(s) is required. The prerequisites for success are perfect sequence identity between at least some of the 21–25 bp siRNAs derived from the dsRNA and the cognate mRNA of the insect pests; and sufficient sequence divergence between all the siRNAs and protein-coding genes of non-target organisms. These analyses can be conducted in silico, by comparing a 21–25 bp moving window along the candidate dsRNA sequence to both the target gene in all target insect taxa, and to all predicted protein-coding genes in all other publicly-available genomes. It may be appropriate to obtain genomic or transcriptomic data for other non-target taxa that currently lack genomic resources, so that the in silico analysis of the proposed dsRNA sequences includes ecologically-relevant organisms. Any proposed dsRNA that fails to yield multiple siRNAs with perfect match to the sequence in all pest insects, or that yields a single siRNA that matches the sequence in any relevant non-target organism should be discarded. Less certain is the degree of sequence mismatch between dsRNA-derived siRNAs and a non-target organism that can be tolerated. Because siRNA molecules can inhibit translation of transcripts with less than perfect sequence identity, the threshold for concern about non-target effects could be less than 100% sequence identity. Further work is required to determine the relative amount of mismatch between the target and effector that causes lack of efficacy. Such results would inform our understanding of how to optimize pest management while minimizing effects to non-target organisms and slowing the evolution of resistance (Section 4.3).

The second issue important for the success of RNAi is delivery at an effective dose, while maintaining acceptable production costs. Recent breakthroughs in dsRNA production methods, which can produce kilogram quantities, continues to reduce the cost associated with dsRNA production and makes it feasible to start discussing strategies which will apply dsRNA products as baits, sprays, or through irrigation systems (Hunter et al., 2010, 2012). In planta RNAi has great potential not only against chewing insect pests [such as the WCRW and CBW studied by Baum et al. (2007) and Mao et al. (2007)], but also against plant sap feeding pests. Transgenic technologies involving expression of toxins from Bacillus thuringiensis (Bt) in crop plants have, contributed little to the control of sucking insect pests, because Bt endotoxins have yet to be identified with activity against these pests (Li et al., 2011). In addition, these insects are becoming increasingly prevalent in Bt crops, as a result of ecological release due to reduced use of broad-spectrum insecticide treatments previously used to control lepidopteran and coleopteran pests (Faria et al., 2007; Lu et al., 2010). Pyramiding RNAi technologies against sap feeders with Bt (or other technologies) against chewing insects could resolve these difficulties. The experimental demonstrations of in planta RNAi against the rice plant hopper Nilaparvata lugens (Zha et al., 2011) and the aphid Myzus persicae (Pitino et al., 2011) provide a proof of principle for this technology.

Alternative approaches are being developed for RNAi delivery as a conventional pesticide, for example as insecticidal baits for urban pests, such as ants, cockroaches and termites (Zhou et al., 2008), or for the aquatic larval stages of mosquitoes (see below). The commercial potential of these methods depends critically on the ability to deliver dsRNA to the target insect, which is in part determined by stability of the dsRNA in the environment, its concentration in the baits and take-up rates by the insects, as offset against the production costs for dsRNA. These objectives will be facilitated by formulations that enhance the uptake of dsRNA into insect cells and its protection against insect dsRNases. For example, dsRNA forms stable 100–400 nm particles in association with chitosan, through the electrostatic forces between the positive charges of the amino group in chitosan and the negatively-charged RNA (McCarroll and Kav pallaris, 2012). Zhang et al. (2010) used the chitosan nanoparticle-based RNAi technology to suppress the expression of two chinin synthase genes (AgCHS1 and AgCHS2) in African malaria mosquito (An. gambiense) larvae. Although this treatment did not kill the larvae, it did reduce the larval chitin content and increased larval susceptibility to the insecticide diflubenzuron.
4.2. RNAi and the protection of insects against parasites and pathogens

The susceptibility of many eukaryotic parasites to RNAi offers a novel strategy to enhance the health of beneficial insects. Of course, this strategy does not apply to bacterial pathogens or the various eukaryotes (e.g. trypanosomes and *Plasmodium* species) which lack any known capacity for RNAi. The opportunity is vividly illustrated by the microsporidian parasite *Nosema* of the honey bee. *Nosema* causes high morbidity and mortality of honey bees (Martín-Hernandez et al., 2011). Two aspects of the biology of *Nosema* make it an especially suitable target for RNAi-based strategies: it has the molecular machinery for RNAi, and it colonizes midgut epithelial cells, a site readily accessed by ingested dsRNA.

When fed honey infected with *N. ceranae* plus dsRNA specific to the *Nosema* ADP/ATP transporter gene, which is essential for *Nosema* energy metabolism, honey bees had a reduced *Nosema* load and lower mortality, together with suppressed transcript abundance of the target genes (Paldi et al., 2010).

Eukaryotic parasites that exploit insect organs other than the gut would be susceptible to RNAi only where the insect host displays systemic spread of the RNAi signal. This has been demonstrated for the ectoparasitic mite, *Varroa destructor*, which feeds on the blood of honey bees (Garbian et al., 2012). When bees were fed on dsRNA specific to a panel of *Varroa* genes, the density of *Varroa* mites on the bees was reduced by up to 50%, with no apparent deleterious effect on the honey bees. The pattern of spread of the RNAi was tested by allowing honey bees to feed on sucrose solution containing dsRNA-GFP (green fluorescent protein; because the genomes of both the insect and mite lack the GFP gene, the distribution of GFP-RNA could be monitored without interference from sequence of endogenous origin). When *Varroa*-infested bees were fed on the test solution, the GFP-RNA was recovered in the *Varroa*. Moreover, when these *Varroa* were subsequently transferred to bees feeding on sugar solution without dsRNA-GFP, the recipient bees acquired the GFP-RNA. These experiments demonstrate that the RNAi can be amplified and spread not only at the level of the individual insect, but also at the colony level in honey bees. Further research is required to establish the frequency and dose of RNAi applications required to sustain protection of colonies, and whether this approach offers a cost-effective strategy for the control of *Varroa* mite, which is of first-order importance in compromising the health of honey bee colonies.

RNAi also holds potential to clear insect vector species from parasites, that themselves are not susceptible to RNAi. A prime example is parasites of the genus *Plasmodium*, the causative agent of malaria (Baum et al., 2009). Conceptually, depletion of proteins required for parasite entry or survival within the insect vector by means of RNAi could be used to create refractory mosquitoites (Brown and Catteruccia, 2006). Proof-of-principle successes have been achieved in the laboratory (Dong et al., 2011). Effective RNAi delivery methodologies that are field-deployable involve oral exposure or transgenic population replacement strategies, and are currently under development.

The natural function of RNAi is protection against viruses, and RNAi has enormous potential in anti-viral therapy. There are opportunities for RNAi-mediated suppression of viral infections in insects, including vectors of socio-economically important viral diseases of humans, livestock and crop plants. Exogenously-applied or ingested dsRNA can be considered as a boost to the native RNAi machinery of the host, conferring protection both by prophylaxis and direct treatment. The value of such boosting is illustrated by research on the titer of various mosquito-vectored arboviruses.

RNAi-mediated antiviral immunity contributes to the suppression of viruses, including dengue virus and Sindbis virus, in the mosquito *Ae. aegypti*, as demonstrated by the increased titer and transmission of these viruses in mosquitoes in which the RNAi machinery was experimentally silenced (Franz et al., 2006; Campbell et al., 2008; Khoo et al., 2010). Furthermore, viral suppression is promoted by enhancing the RNAi pathway, achieved by engineering the insects to express an inverted-repeat RNA that triggers production of dsRNA specific to the virus sequence (Franz et al., 2011; Mathur et al., 2010). There is some evidence for viral-mediated suppression of RNAi, for example by the Sindbis virus and West Nile virus, in mosquito cells (Crimotitch et al., 2009; Schnettler et al., 2012). Such suppression may be the reason why supplementary dsRNA is required to achieve RNAi-mediated elimination of viral infection from the insect host.

Another insect system demanding urgent solutions to viral infections is the honey bee, especially in the context of evidence that viruses, including the Israeli acute paralysis virus (IAPV), may contribute to the ongoing decline of honey bees, including colony collapse disorder (Evans and Schwarz, 2011). Evidence that exogenous dsRNA can supplement the endogenous RNAi machinery comes from the demonstration that IAPV infection of honey bees can be eliminated by orally-delivered dsRNA corresponding to two different sequences of the IAPV genome (Maori et al., 2009). Among colonies inoculated with IAPV, mortality was reduced in those treated with IAPV-dsRNA relative to those that were not treated or that were treated with non-IAPV dsRNA. These results led to large-scale field test in the USA, in which honey bees were fed a dsRNA product, Remebee-I, in the presence of the IAPV (Hunter et al., 2010). Honey bee survival, colony size and honey production were all increased in the Remebee-I treatment. Ingested IAPV-specific dsRNA successfully reduced the negative effects of IAPV infection in 160 honey bee hives in two states (Florida and Pennsylvania) with very different climates and seasons. These results provide the first successful field demonstration of the use of RNAi as a large scale preventative treatment for an insect disease.

The antiviral effect of RNAi has also been successfully augmented for disease control in a non-insect arthropod, the cultured shrimp, *L. vannamei*. Diseases caused by viruses are economically devastating to the shrimp industry, and induced RNAi provides protection from a number of different viruses, including single- and double-stranded RNA viruses and a DNA virus (Bartholomay et al., 2012). This strategy enhances RNAi-based antiviral immunity, providing long-term, highly specific protection and a route for vaccination of cultured shrimp against viral diseases. For example, dsRNA designed to target the 5’ end of ORF1 in the genome of Infectious myonecrosis virus (IMNV) provides significant disease protection even 52 days after vaccination (Loy et al., 2012). Moreover, the same RNAi trigger was used to provide therapeutic effect such that disease pathology resolved and 50% of animals survived if the RNAi trigger was provided within 48 h post-infection (Loy et al., 2012). The outstanding challenge is a viable delivery strategy, because shrimp culture involves hundreds of thousands of animals in hectare-sized ponds.

4.3. The evolutionary stability of RNAi-based management of insect populations

The relationship between viruses and RNAi-based insect immunity is evolutionarily dynamic. This is indicated by both the presence of viral suppressors of RNAi (see above) and the positive selection on the genes contributing to RNAi-machinery interacting with siRNAs, but not the endogenous miRNAs (Obbard et al., 2006). We can, therefore, anticipate that insects, viruses and eukaryotic parasites will respond to strong selection exerted by RNAi-based control strategies. For example, insects that carry viruses with RNAi suppressors would be at a selective advantage on RNAi-protected crops, and RNAi-based prophylactics for honey bee colonies would select for viral pathogens with RNAi suppression. The RNAi
suppression mechanisms that have evolved are not specific to a particular target sequence. This implies that resistance to a dsRNA specific to one gene cannot be prevented by pyramiding multiple genes with different function, nor overcome by switching to a different gene or gene set.

The genetic variation that exists within and among insect populations could also present a challenge to the application of RNAi for pest control, depending on the amount of mismatch present between the dsRNA and the target transcript. Furthermore, single nucleotide polymorphisms (SNPs) that result in lower effectiveness of the RNAi, could potentially be selected for and lead to the evolution of resistance. If such SNPs were synonymous they would be expected to have little or no fitness cost in the absence of the selecting agent (dsRNA), and resistance could evolve rapidly. However, the degree of mismatch (i.e. the number of SNPs) that would be needed to prevent RNAi from controlling a pest is not known. The long-term benefits of RNAi-based applications in insect pest management will require new and independent thought on effective resistance management strategies designed to minimize selective pressures and delay the evolution of resistance.

4.4. RNAi risks and regulation

The above examples offer clear evidence for potential applications for RNAi for the control of insect pests, manipulation of insect disease vectors, and management of beneficial insects, together with concerns about the stability of RNAi strategies in the face of selection for resistance. Overlying these considerations is a very real uncertainty regarding the environmental and ecological risks posed by these technologies. The Federal regulatory framework for estimating the ecological risks associated with RNAi technologies is still in development, and a number of critical gaps remain including potential toxicity to non-target organisms (see Section 4.1), environmental fate, and importantly, the risk of resistance evolution in target pests (Section 4.3). Documenting efficacy of the technology is ongoing and regulatory considerations for RNAi-based insecticidal traits, such as the development of standardized environmental risk assessment, are still being developed (Auer and Frederick, 2009; http://cera-gmc.org/docs/cera_publications/pub_08_2011.pdf). Considerations of how to evaluate sequence specificity, environmental fate, and exposure of non-target organisms are still being developed. However, US regulatory agencies such as the Environmental Protection Agency and the Department of Agriculture have provided preliminary assessments (http://cera-gmc.org/docs/cera_publications/pub_08_2011.pdf) suggesting that data requirements for RNAi traits may be reduced based primarily on the lack of a plant incorporated protein, such as a Bt toxin. There is also a lack of information on the risk of insect resistance to RNAi-mediated control that is a critical impediment to the development of an insect resistance management plan aimed at promoting a responsible and sustainable use of the technology. Insecticide resistance presents a major challenge for the sustainable control of pests. In the case of insects, pest species have found ways to evolve resistance to nearly every control strategy that has been used. Predictions that resistance could not develop to a new control strategy (e.g. Williams, 1967) have proven to be wrong time and time again.

5. Concluding comments

A decade of research on RNAi in insects has demonstrated the great power of the technology for discovery-led science and potential for improved management of insect populations. As the science has matured, it has equally become evident that RNAi is no panacea, but introduces a range of new conceptual and technological challenges for insect scientists. Insects vary widely in their amenability to RNAi, and no single protocol is suitable for all species. Against the backdrop of this functional diversity, it is unfortunate that there has been a dearth of systematic investigation of the mechanisms of RNAi in insects. We still have only a weak understanding of whether and how the RNAi signal is amplified in individual cells and disseminated between cells in insects. It is increasingly recognized that the caveats in our understanding of insect-specific mechanism are a major limitation to the implementation of RNAi. A priority for the future is for the insect research community to apply their persistence and ingenuity to solve the fundamentals of how insect RNAi works, in the context of the physiology of the insect body, and apply that to the pressing problems posed by pests and beneficial insects.

Acknowledgements

We thank all of the speakers and participants at the 2012 ESA symposium “RNAi: The Power, the Promise and the Frustration”, from which the idea for this paper originated. This work was supported by the following grants: NIH R01 AI095842 to K.M., NSF 1114370 to L.C.B., ACSF to J.G.S. and AFRI-NIFA NYW-2011-04650 to A.E.D.

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