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研究課題名(和文)Origins, spread and evolution of novel honey bee parasites and diseases

研究課題名(英文)Origins, spread and evolution of novel honey bee parasites and diseases

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研究成果の概要(和文):寄生生物によるホストスイッチの成功は、新たな資源を獲得し種分化を起こす進化上の出来事である。観察は困難で世界的なパンデミックの要因である。我々は、外部寄生虫として世界的にミッバチ減少をもたらしているVarroa属の二種のダニについて、ホストスイッチの動態の変遷を再現した。このダニがクローン様に均質とする従来の考えと、世界的に広まる現状には遺伝学的なある。我々は数百個体に相当する遺伝的多様性がホストスイッチに関与することを発見し、ダニが多様なミッバチ集団や殺虫剤に適応する根拠を見出した。これにより、ダニの動態に関わる変数のうち特に遺伝的多様性がホストスイッチに重要なことが明らかとなった。

研究成果の学術的意義や社会的意義

Our results highlight the importance of demographic parameters, particularly genetic diversity, for parasitic host switches

研究成果の概要(英文): Successful host switches by parasites are evolutionary events that enable them to access new resources, and can lead to eventual speciation. Host switch events can be difficult to observe, yet they can lead to worldwide pandemics. We reconstructed the demographic history of host switches by two species of ectoparasites, Varroa mites, which are the principal drivers of worldwide honey bee declines. Varroa mites are believed to be "quasi-clonal", making their worldwide success a genetic paradox. However, we found that the host switch involved the genetic equivalent of hundreds of individuals, explaining the remarkable adaptation of the parasites to both diverse bee populations and pesticides. These results highlight the importance of demographic parameters, particularly genetic diversity, for parasitic host switches.

研究分野: evolution

キーワード: evolution parasites honey bees

1.研究開始当初の背景

Arms races between parasites and their hosts drive evolutionary innovation. Novel parasites can decimate host populations or drive them extinct, unless counter-adaptations evolve. Similarly, parasite evolution accompanies the very act of host switching, since it requires adaptations to novel host physiology in order to persist and to spread. Because parasite adaptations tend to be host-specific, host switches are often associated with host-associated genetic differentiation and eventual speciation. However, only the endpoint of this process is typically observed, as host switches tend to occur rapidly, and the original host is often unknown. As a result, many unanswered questions remain about how parasites acquire new hosts. For instance, if host switches are accompanied by a bottleneck due to reduced gene flow from the ancestral host, how does the parasite have sufficient genetic diversity to adapt? Does gene flow cease completely, or does it continue at a low level, potentially providing additional genetic material for adaptations?

2.研究の目的

One of the major limiting factors for host switching is the geographic separation between parasites and potential hosts. Globalization has eased these barriers, sometimes giving rise to pandemics (Hatcher, Dick, and Dunn 2012). As a result, host switches are easier to observe and to study in something approaching real time. One of the most dramatic and economically important switches involved the two ectoparasitic mite species, *Varroa destructor* and *Varroa jacobsoni*, which acquired the western honey bee (*Apis mellifera*) as a new host, ~70 and ~12 years ago, respectively (Anderson and Trueman 2000; Roberts, Anderson, and Tay 2015). *V. destructor*, in particular, spread worldwide, causing extensive honey bee population collapses, whereas *V. jacobsoni* has so far remained in Oceania (Traynor et al. 2020; Noël, Le Conte, and Mondet 2020). Both mites were originally found on the sister species, *Apis cerana*, and came into contact with *A. mellifera*, which was brought in for purposes of beekeeping (Crane 1968; Chantawannakul et al. 2016). These two host switches occurred in parallel and relatively recently, allowing the reconstruction of how the host switches took place, using genomic tools.

Both switches have been investigated using microsatellite markers and relatively short mitochondrial markers, which revealed that in both species, populations parasitizing A. mellifera were strongly differentiated and genetically depauperate (Solignac et al. 2005; Warrit, Smith, and Lekprayoon 2006; Navajas et al. 2010; Beaurepaire et al. 2015; Roberts, Anderson, and Tay 2015; Dietemann et al. 2019). While V. destructor, in particular, was described as "quasi-clonal" (Solignac et al. 2005), it successfully spread worldwide, and has shown a remarkable ability to parasitize genetically diverse A. mellifera populations, as well as to evolve resistance to human counter-measures, such as pesticides (González-Cabrera et al. 2016). How does a bottlenecked species achieve such a level of success? Increasing evidence from population genomic analysis of fungal pathogens suggests that the success of many pathogens appears to rely on maintaining some level of adaptive diversity despite the presence of bottlenecks during host switches (Fry 2016; McMullan et al. 2018), but similar work is lacking in animal systems.

3.研究の方法

Using whole-genome sequencing on 63 mites collected in their native ranges from both the ancestral and novel hosts, we were able to reconstruct the known temporal dynamics of the switch using a combination of whole genome sequencing, both nuclear and mitochondrial.

4. 研究成果

In order to answer this question and to gain broader insight into how host switches happen, we sequenced genomes from sympatric populations of the two mites across Asia

and Oceania, collected on both novel and introduced hosts. This allowed us far greater power to examine how the host switch took place with much greater precision than was possible previously. We found strikingly parallel dynamics at play in both host switches, which were characterized by a surprisingly large effective population size at the time of the switch and ongoing gene flow with cryptic population genetic processes that may have helped *Varroa* succeed.

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5 . 主な発表論文等

〔雑誌論文〕 計2件(うち査読付論文 2件/うち国際共著 2件/うちオープンアクセス 2件)

1.著者名	4 . 巻
Maeva A. Techer, Rahul V. Rane, Miguel L. Grau, John M.K. Roberts, Shawn T. Sullivan,Ivan	2
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1.発表者名

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3 . 学会等名

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2018年

〔図書〕 計0件

〔産業財産権〕

〔その他〕

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6.研究組織

0	. 加力光組織		
	氏名 (ローマ字氏名) (研究者番号)	所属研究機関・部局・職 (機関番号)	備考

7. 科研費を使用して開催した国際研究集会

〔国際研究集会〕 計0件

8. 本研究に関連して実施した国際共同研究の実施状況

共同研究相手国	相手方研究機関
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