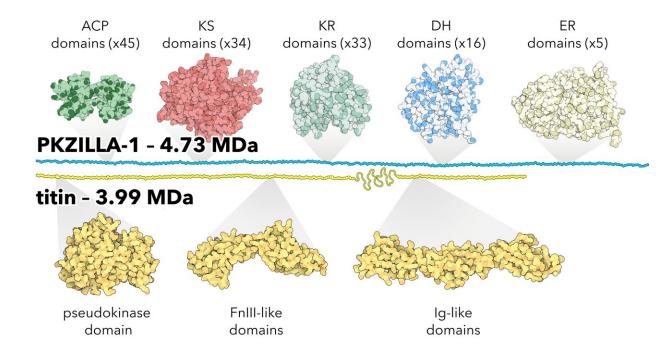


Largest protein yet discovered builds algal toxins

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Stylistic size comparison of PKZILLA-1 and human titin, not to scale, wherein PKZILLA-1 is shown as roughly 25% larger than titin. Abbreviations: megadalton (MDa), acyl carrier protein (ACP), ketosynthase (KS), ketoreductase (KR), dehydratase (DH), enoylreductase (ER), fibronectin (Fn), immunoglobulin (Ig). Titin figure adapted from https://pdb101.rcsb.org/motm/185 under Creative Commons license, attribution: David S. Goodsell, RCSB Protein Data Bank (PDB). ACP structure modeled from bacterial ACP (PDB id: 2FAE). KS, KR, DH, ER, modeled from mammalian fatty acid synthase (PDB id: 2VZ8). Credit: Tim Fallon, Ph.D.



While seeking to unravel how marine algae create their chemically complex toxins, scientists at UC San Diego's Scripps Institution of Oceanography have discovered the largest protein yet identified in biology.

Uncovering the biological machinery the algae evolved to make its intricate <u>toxin</u> also revealed previously unknown strategies for assembling chemicals, which could unlock the development of new medicines and materials.

Researchers found the <u>protein</u>, which they named PKZILLA-1, while studying how a type of algae called Prymnesium parvum makes its toxin, which is responsible for massive fish kills.

"This is the Mount Everest of proteins," said Bradley Moore, a marine chemist with joint appointments at Scripps Oceanography and Skaggs School of Pharmacy and Pharmaceutical Sciences and senior author of a new study detailing the findings. "This expands our sense of what biology is capable of."

PKZILLA-1 is 25% larger than titin, the previous record holder, which is found in human muscles and can reach 1 micron in length (0.0001 centimeter or 0.00004 inch).

Published in <u>Science</u> the study shows that this giant protein and another super-sized but not record-breaking protein—PKZILLA-2—are key to producing prymnesin—the big, complex molecule that is the algae's toxin.

In addition to identifying the massive proteins behind prymnesin, the study also uncovered unusually large genes that provide Prymnesium parvum with the blueprint for making the proteins.



Finding the genes that undergird the production of the prymnesin toxin could improve monitoring efforts for <u>harmful algal blooms</u> from this species by facilitating water testing that looks for the genes rather than the toxins themselves.

"Monitoring for the genes instead of the toxin could allow us to catch blooms before they start instead of only being able to identify them once the toxins are circulating," said Timothy Fallon, a postdoctoral researcher in Moore's lab at Scripps and co-first author of the paper.

Discovering the PKZILLA-1 and PKZILLA-2 proteins also lays bare the alga's elaborate cellular assembly line for building the toxins, which have unique and complex chemical structures. This improved understanding of how these toxins are made could prove useful for scientists trying to synthesize new compounds for medical or industrial applications.





Golden alga (Prymnesium parvum) fish kill - Lake Granbury March 2007. Credit: Texas Parks and Wildlife Department/TPWD

"Understanding how nature has evolved its chemical wizardry gives us as scientific practitioners the ability to apply those insights to creating useful products, whether it's a new anti-cancer drug or a new fabric," said Moore.

Prymnesium parvum, commonly known as golden algae, is an aquatic single-celled organism found all over the world in both fresh and saltwater. Blooms of golden algae are associated with fish die offs due to its toxin prymnesin, which damages the gills of fish and other water



breathing animals.

In 2022, a golden algae bloom killed <u>500-1,000 tons of fish in the Oder River</u> adjoining Poland and Germany. The microorganism can cause havoc in aquaculture systems in places ranging from Texas to Scandinavia.

Prymnesin belongs to a group of toxins called polyketide polyethers that includes brevetoxin B, a major red tide toxin that regularly impacts Florida, and ciguatoxin, which contaminates reef fish across the South Pacific and Caribbean. These toxins are among the largest and most intricate chemicals in all of biology, and researchers have struggled for decades to figure out exactly how microorganisms produce such large, complex molecules.

Beginning in 2019, Moore, Fallon and Vikram Shende, a postdoctoral researcher in Moore's lab at Scripps and co-first author of the paper, began trying to figure out how golden algae make their toxin prymnesin on a biochemical and genetic level.

The study authors began by sequencing the golden alga's genome and looking for the genes involved in producing prymnesin. Traditional methods of searching the genome didn't yield results, so the team pivoted to alternate methods of genetic sleuthing that were more adept at finding super long genes.

"We were able to locate the genes, and it turned out that to make giant toxic molecules this algae uses giant genes," said Shende.

With the PKZILLA-1 and PKZILLA-2 genes located, the team needed to investigate what the genes made to tie them to the production of the toxin. Fallon said the team was able to read the genes' coding regions like sheet music and translate them into the sequence of amino acids that



formed the protein.

When the researchers completed this assembly of the PKZILLA proteins they were astonished at their size. The PKZILLA-1 protein tallied a record-breaking mass of 4.7 megadaltons, while PKZILLA-2 was also extremely large at 3.2 megadaltons. Titin, the previous record-holder, can be up to 3.7 megadaltons—about 90-times larger than a typical protein.





Prymnesium parvum cell. Credit: Greg Southard, Texas Parks and Wildlife Department

After additional tests showed that golden algae actually produce these giant proteins in life, the team sought to find out if the proteins were



involved in making the toxin prymnesin. The PKZILLA proteins are technically enzymes, meaning they kick off <u>chemical reactions</u>, and the team played out the lengthy sequence of 239 chemical reactions entailed by the two enzymes with pens and notepads.

"The end result matched perfectly with the structure of prymnesin," said Shende.

Following the cascade of reactions that golden algae uses to make its toxin revealed previously unknown strategies for making chemicals in nature, said Moore. "The hope is that we can use this knowledge of how nature makes these complex chemicals to open up new chemical possibilities in the lab for the medicines and materials of tomorrow," he added.

Finding the genes behind the prymnesin toxin could allow for more cost effective monitoring for golden algae blooms. Such monitoring could use tests to detect the PKZILLA genes in the environment, akin to the PCR tests that became familiar during the COVID-19 pandemic. Improved monitoring could boost preparedness and allow for more detailed study of the conditions that make blooms more likely to occur.

Fallon said the PKZILLA genes the team discovered are the first genes ever causally linked to the production of any marine toxin in the polyether group that prymnesin is part of.

Next, the researchers hope to apply the non-standard screening techniques they used to find the PKZILLA genes to other species that produce polyether toxins.

If they can find the genes behind other polyether toxins, such as ciguatoxin, which may affect <u>up to 500,000 people annually</u>, it would open up the same genetic monitoring possibilities for a suite of other



toxic algal blooms with significant global impacts.

More information: Timothy R. Fallon et al, Giant polyketide synthase enzymes in the biosynthesis of giant marine polyether toxins, *Science* (2024). DOI: 10.1126/science.ado3290. www.science.org/doi/10.1126/science.ado3290

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