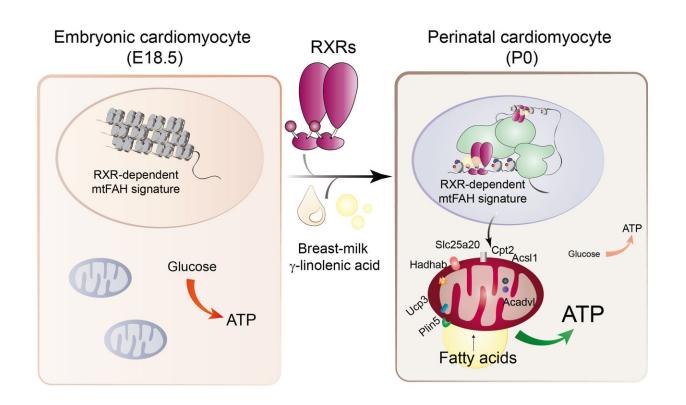


GLA, the fatty acid that helps the heart to function properly after birth

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After birth, GLA activates RXR in the postnatal cardiomyocytes. This results in the activation of genetic programs that ensure that the cardiomyocyte mitochondria mature and use lipids as the energy source for the generation of ATP. This metabolic adaptation makes sure that the myocardium is provided with enough energy to sustain the heartbeat outside the womb, thus ensuring postnatal survival. Credit: CNIC

A study conducted in mice and led by researchers at the Centro Nacional



de Investigaciones Cardiovasculares (CNIC) has revealed that maternal milk provides an essential signal that triggers the maturation of heart metabolism after birth, allowing the neonatal heart to function correctly and ensuring postnatal survival.

The study shows that the fatty acid gamma-linolenic acid (GLA), present in <u>breast milk</u>, binds to the retinoid X receptor (RXR) protein found in <u>heart cells</u>. RXR acts as a nutritional sensor of lipids and vitamin A derivatives, altering <u>gene expression</u> and influencing <u>biological functions</u> such as immunity, cell differentiation, and metabolism. Once activated by maternal GLA, RXR initiates genetic programs that equip mitochondria, the energy centers of the cell, with the enzymes and other proteins they need to to start consuming lipids, the primary source of energy in the mature <u>heart</u>.

The results, published May 24, 2023 in *Nature*, could have significant therapeutic implications for cardiovascular disorders involving mitochondrial and metabolic dysfunction, as well as diseases related to alterations in postnatal developmental processes, explained study leader Dr. Mercedes Ricote, who heads the Nuclear Receptor Signaling group at the CNIC.

In a <u>mouse model</u>, the research team found that the absence of RXR in the heart or the lack of GLA in maternal milk prevented mitochondria in the hearts of newborn mice from producing energy correctly, leading to severe heart failure and death 24 to 48 hours after birth.

The heart of a newborn mammal must quickly produce energy to sustain cardiac contraction outside of the womb. To achieve this, cardiomyocytes, the contracting cells of the myocardium (the cardiac muscle), need to activate their mitochondria to use lipids as the energy source for the generation of ATP (adenosine triphosphate—the essential energy currency of the cell). Although this process is crucial for the



survival of the organism, until now, little was known about the signals that trigger the physiological adaptation of the heart after birth.

"The need to maintain a constant and uninterrupted beat places an immense energy demand on the heart," explained Dr. Ricote. "To meet their energy needs, cardiomyocytes maintain a tight control over the cellular pathways that produce energy. Any imbalance in these bioenergetic mechanisms can lead to the development of serious cardiovascular pathologies."

For Dr. Ricote, part of the study's novelty "lies in demonstrating that RXR plays a critical role in cardiac muscle, contrary to what was previously thought. This is an important conceptual advance in the field of nuclear receptors."

According to first author Dr. Ana Paredes, the study presents a new framework for understanding the postnatal adaptations that occur in newborn mammals to meet the requirements of the extrauterine environment. "Birth is a physiological challenge for the newborn," Dr. Paredes explained. "With this study, we show that <u>maternal milk</u>, besides its nutritional function, plays a signaling role by instructing cardiomyocytes that they need to activate their metabolism because they are no longer supported by maternal physiology."

In conclusion, the study shows that the fatty acid GLA, found in breast milk, is the key signal that ensures correct cardiac function after birth. GLA activates the cell protein RXR, which then directs coordinated gene expression changes to ensure that cardiomyocyte mitochondria mature so that they can produce energy in the extrauterine environment.

The results open the way to treatments to modulate RXR activity in cardiomyocytes with specific drugs, including some that already have approval from the US Federal Drug Administration for cancer



treatments. "Our study proposes RXR as a possible therapeutic target for neonatal heart disorders and systemic diseases triggered by metabolic errors," concluded Dr. Ricote.

The research involved multidisciplinary approaches and advanced sequencing techniques, including key contributions from the CNIC teams led by Dr. José Antonio Enríquez, Dr. Fátima Sánchez-Cabo, and Dr. Jesús Vázquez.

More information: Mercedes Ricote, γ-Linolenic acid in maternal milk drives cardiac metabolic maturation, *Nature* (2023). DOI: 10.1038/s41586-023-06068-7. www.nature.com/articles/s41586-023-06068-7

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