

IISER Mohali uncovered how *Salmonella* survives in human cells

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A team of scientist at the Institute of Science and Research (IISER), Mohali has uncovered mechanism of how *Salmonella* manages access to membranes and nutrients in human cells through structures (known as lysosomes) that hold a variety of proteins for its own growth and survival.

The research was led by Dr Mahak Sharma along with other team members including Aastha Sindhwani, Subhash B. Arya, Harmeet Kaur, Divya Jagga, Amit Tuli, Mahak Sharma. The results of the study have been published in journal PLOS Pathogens.

Generally, many disease-causing bacteria are insidious. They try to manipulate intracellular structures of human cells for their own growth and survival. Among the bacterial pathogens which use human proteins for thriving are *Mycobacterium tuberculosis* which causes tuberculosis and *Salmonella enterica typhimurium*, causative agent of typhoid and gastroenteritis.

According to the study, pathogens have evolved several tactics to manipulate the activity of host proteins, all aimed at creating their intracellular replicative niche (known as a vacuole). For example, *M. tuberculosis* sucks iron and other nutrients from the host cells to prevent degradation of its own niche. On the other hand, *Salmonella* acquires both membrane and nutrients within a mature acidic vacuole. It avoids bacteria-killing enzymes by rerouting them out of the host cell as well as by forming an extensive network of membranes with its vacuole that dilutes these enzymes.

“We are trying to understand how lysosomal proteins; specifically the small GTP-binding proteins and their effectors facilitate cargo delivery to lysosomes. Previous studies have shown that *Salmonella* vacuole acquires several characteristics of the host cell lysosomes for nutrient access from this organelle but the mechanism at play was not known,” Dr. Mahak Sharma told India Science Wire.

Dr. Amit Tuli of CSIR-IMTECH, Chandigarh, who collaborated for the study, further explained that “Salmonella-containing vacuole (SCV) does not inhibit maturation of its vacuole but rather acquires several features of host lysosomes. Indeed, acidification of the vacuole is required for production of virulence factors that in turn help Salmonella to acquire both membranes and nutrients from the host endosomes and lysosomes.”

Researchers say Salmonella secretes a protein known as SifA that recruits a host chemical, HOPS complex, to SCV membranes. Recruitment of HOPS complex is key step leading to docking and fusion of SCVs with host endosomes and lysosomes. Blocking HOPS function prevents Salmonella replication inside its vacuolar niche but cutting off nutrient supply.

"Based on our findings, we envision that small molecules or peptide-based inhibitors that specifically block the interaction of Salmonella protein SifA with host factors, including HOPS complex, can potentially prevent Salmonella replication and infection in the human host," added Dr Sharma.