Malate synthase contributes to the survival of *Salmonella* Typhimurium against nutrient and oxidative stress conditions

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To survive and replicate in the host, *S.* Typhimurium have evolved several metabolic pathways. The glyoxylate shunt is one such pathway that can utilize acetate for the synthesis of glucose and other biomolecules. This pathway is a bypass of the TCA cycle in which CO₂ generating steps are omitted. Two enzymes involved in the glyoxylate cycle are isocitrate lyase (ICL) and malate synthase (MS). We determined the contribution of MS in the survival of *S.* Typhimurium under carbon limiting and oxidative stress conditions. The *ms* gene deletion strain (∆*ms* strain) grew normally in LB media but failed to grow in M9 minimal media supplemented with acetate as a sole carbon source. However, the ∆*ms* strain showed hypersensitivity (*p* < 0.05) to hypochlorite. Further, ∆*ms* strain has been significantly more susceptible to neutrophils. Interestingly, several folds induction of *ms* gene was observed following incubation of *S.* Typhimurium with neutrophils. Further, ∆*ms* strain showed defective colonization in poultry spleen and liver. In short, our data demonstrate that the MS contributes to the virulence of *S.* Typhimurium by aiding its survival under carbon starvation and oxidative stress conditions.

Based on antigenic presentations¹, *Salmonella enterica* serovars are grouped as typhoidal and non-typhoidal *Salmonella* (NTS). WHO recognizes NTS as one of the three most common food-borne bacterial diseases in humans all over the world. Old, young, and immunocompromised individuals are highly prone to *Salmonella* infection². Among NTS, serovar Typhimurium is most commonly isolated from patients around the globe³.

Following ingestion, a proportion of the microorganisms resists the low gastric pH, invades the intestinal mucosa, and replicates in the sub-mucosa and Peyer’s patches⁴. Following intestinal penetration, *S.* Typhimurium gains access to the mesenteric lymph nodes, where the bacteria are engulfed by phagocytic cells, such as macrophages⁵. Once inside the macrophages, *S.* Typhimurium is compartmentalized into a modified vacuole known as "*Salmonella*-containing vacuole" (SCV) and represents a central feature in the intracellular survival and growth of this bacterium⁶. Thus, the engulfment by the macrophage thrusts the *S.* Typhimurium into an alien milieu which is rich in various antimicrobials and devoid of key nutrients essential for metabolism and replication. To survive under such harsh conditions, *S.* Typhimurium modulates the functions of phagocytes in several ways. First, effectors encoded by the type III secretion system of *S.* Typhimurium impede the assembly of phagosomal oxidase and consequently inhibit the production of superoxide radicals. Second, the SCV acts as a shield for *S.* Typhimurium that not only prevents lysosomal fusion but also limits the exposure of contained bacterial cells to antimicrobial agents⁷. While the primary antioxidants of *S.* Typhimurium directly quench oxidants, the repair enzymes restore the functions of the damaged biomolecules⁸,⁹.

However, survival against the antimicrobial assault in the phagolysosome depends on the microbe’s ability to synthesize the proteins and other biomolecules required to counteract stresses. Thus, a pathogen must find the requisite nutrients to provide the building blocks for these complex macromolecules and the energy with which to synthesize them¹⁰. It is the metabolic flexibility of *S.* Typhimurium which allows it to survive in such harsh conditions within the host¹¹. The ability to fulfill its nutrient requirements from alternate sources might play an important role in the adeptness of *S.* Typhimurium in the host. One such survival mechanism is the existence of the glyoxylate cycle, whose primary function is to permit bacterial/cellular growth when C₂ compounds, such as ethanol and acetate, are the only sources of carbon¹². Few studies suggest that macrophages are rich in fatty
Malate synthase is susceptible to oxidative damage. Earlier, we have reported oxidant mediated oligomerization and Met-SO formation in malate synthase. To corroborate our findings, if malate synthase is prone to oxidation, we assessed the carbonylation by Oxyblot. Carbonylation is a well-accepted marker to assess protein oxidation. Our Oxyblot analysis showed a much more intense band in oxidant exposed malate synthase. Furthermore, Met residues of MS are prone to oxidation, and can be repaired by methionine sulfoxide reductase. However, the role of MS in the survival of S. Typhimurium under oxidative stress conditions is not known. In the present study, we aimed to explore the contribution of ms in the survival of S. Typhimurium during oxidative stress and colonization in the liver and spleen of poultry. By employing various biochemical, molecular biology tools along with cell culture and live animal studies we show that the ms is required for growth of S. Typhimurium under carbon starvation and survival under oxidative stress conditions. Finally, our data suggest that the ms contributes to the survival of S. Typhimurium in phagocytic cells and poultry spleen and liver.

Results
Δms strain does not exhibit defective growth in LB broth. The growth rate of Δms strain was comparable to that of S. Typhimurium in LB broth. However, the Δms strain failed to grow in M9 media supplemented with acetate as the sole carbon source (Fig. 1 and Supplementary Fig. 1). Further, we observed the induction of malate synthase protein in S. Typhimurium cultured in M9 media supplemented with acetate (Fig. 1 and Supplementary Fig. 2). The immunoreactivity of MS band was highly intense in the lysates of S. Typhimurium cultured in the M9 media supplemented with acetate as compared to bacteria cultured in glucose added media (Supplementary Fig. 2a). However, the loading on SDS-gel was almost similar for both the conditions (Supplementary Fig. 2b).

Δms strain shows hypersusceptibility to HOCl. HOCl is one of the most potent oxidants generated by myeloperoxidase catalyzed reaction between H₂O₂ and chloride ions. We assessed the susceptibility of Δms strain to HOCl. The percent recovery of the Δms strain was found to be reduced up to 9.39% and 0.013% as compared to that of 42.31% and 0.141% in the case of WT after treating the cells with 1.5 and 3 mM HOCl respectively (Fig. 2).

Δms strain is highly susceptible to neutrophils. Neutrophils are the key HOCl producing cells. After observing susceptibility of Δms strain to reagent HOCl, we investigated the role of ms gene in the survival of S. Typhimurium against neutrophil-mediated killing. Neutrophils were incubated with WT or Δms strains of S.

The potential contribution of the glyoxylate cycle in the survival of microorganisms under oxidative stress and pathogenesis has been suggested. Few studies in fungal and bacterial pathogens have indicated the upregulation of glyoxylate cycle genes following phagocytosis. An increased metabolic flux (48%) through the glyoxylate shunt is observed in E. coli experiencing superoxide stress. A recent study in M. tuberculosis showed the increased susceptibility of ms knockdown strain to oxidative and nitrosative stresses, and macrophages. It has been suggested that ICL is required for the persistence of Salmonella during chronic infection in mice.

Earlier, we have shown that the ms gene deletion strain showed defective colonization in poultry caecum. Further, Met residues of MS are prone to oxidation and can be repaired by methionine sulfoxide reductase. However, the role of MS in the survival of S. Typhimurium under oxidative stress conditions is not known. In this study, we aimed to explore the contribution of ms in the survival of S. Typhimurium during oxidative stress and colonization in the liver and spleen of poultry. By employing various biochemical, molecular biology tools along with cell culture and live animal studies we show that the ms is required for growth of S. Typhimurium under carbon starvation and survival under oxidative stress conditions. Finally, our data suggest that the ms contributes to the survival of S. Typhimurium in phagocytic cells and poultry spleen and liver.

Figure 1. Δms strain fails to grow on acetate as a sole carbon source: overnight grown cultures of WT and Δms strains of S. Typhimurium were diluted in LB broth (A) or M9 media supplemented with either 0.4% glucose (B) or 0.4% acetate (C) and grown on a shaker incubator. The aliquots were taken at an interval of 1 h and optical densities were measured at 600 nm. Data is presented as mean ± SD (n = 3).
Typhimurium. By retrospective plating, the observed MOI for WT and Δms strains were 1:7 and 1:8.5 (neutrophil: bacteria), respectively. Following 15 min of incubation, the percent recovery of the Δms strain was found to be reduced up to 8.47% as compared to that 16.42% in case of WT after incubation with neutrophils (Fig. 3). Indeed, the Δms strain was significantly more susceptible (p < 0.0001) to neutrophil mediated killing.

Incubation of S. Typhimurium with neutrophils induces expression of malate synthase. After observing the hypersusceptibility of Δms strain to neutrophils, we evaluated the expression of ms following incubation of S. Typhimurium with neutrophils. The S. Typhimurium was co-cultured with neutrophils for 15 min and the relative expression of ms gene was analyzed by qRT-PCR. We observed 3.82 folds upregulation of ms in S. Typhimurium following incubation with neutrophils (Supplementary Fig. 6).

Δms strain is attenuated in spleen and liver. Poultry is one of the most important reservoirs of S. Typhimurium in nature which transmits infection through eggs and meat. After a short enteric phase, S. Typhimurium enters the spleen and liver where it targets various phagocytic cells. We determined the bacterial loads in the poultry spleen and liver on 7, 14 and 21 days post-infection. In the spleen of WT strain infected birds, we obtained bacteria at all times post-infection. The numbers of S. Typhimurium recovered on 7, 14 and 21 days were expressed in log10 CFU/spleen (mean ± SE). Number of WT bacteria recovered were 1.4 ± 0.36, 1.3 ± 0.61 and 0.6 ± 0.37 respectively. However, the survival of Δms strain was compromised in the spleen and we recovered mutant bacteria only up to 14 days post-infection. The counts were 0.58 ± 0.36 and 0.64 ± 0.4 on days 7 and 14th respectively (Fig. 4).

In the liver, we recovered bacteria from both WT and Δms strain infected birds until 14 days post-infection and expressed the values as log10 CFU/ gm of the liver (mean ± SE). The numbers of WT bacteria recovered were 0.86 ± 0.5 and 2.37 ± 1.0 on 7 and 14 days post-infection, respectively. In Δms strain infected birds, we recovered
0.4 ± 0.4 and 0.4 ± 0.4 on 7th and 14th post-infection, respectively (Fig. 5). On 14 days post-infection the bacte-
rial loads in the liver of ∆ms strain infected birds were significantly (p < 0.05) lower than the WT-infected group.

Discussion

To thrive within the host, S. Typhimurium has to confront nutrient/ carbon limiting and oxidant rich environ-
ment of the phagocytic cells. To assess the role of the glyoxylate cycle in S. Typhimurium under these two (carbon
starvation and oxidative) stress conditions, we deleted the malate synthase gene and confirmed the deletion by
PCR9. Δms strain grew normally in LB media or M9 minimal media supplemented with glucose (Fig. 1A,B and
Supplementary Fig. 1). However, Δms strain displayed total growth defect when cultured in M9 minimal media
supplemented with acetate as a sole carbon source (Fig. 1C and Supplementary Fig. 1).

Induction of glyoxylate shunt enzymes has been reported in S. Typhimurium cultured in acetate containing
media22. We observed huge (more than 20 folds) induction of malate synthase protein in S. Typhimurium grown
in M9 minimal media supplemented with acetate as compared to cultured in M9 minimal media supplemented
with glucose (Supplementary Fig. 2). These data suggest a pivotal role of malate synthase in the survival of S.
Typhimurium under carbon limiting conditions.

Under oxidative stress, the activation of the enzymes of the glyoxylate cycle has been observed in vari-
ous microorganisms including, Pseudomonas aeruginosa23, M. tuberculosis24, Alishewanella25 and Acinetobacter
oleivorans26. Bioinformatics analysis suggests that the microorganisms in which the glyoxylate cycle is func-
tional, are either aerobic or facultative anaerobic23. The use of oxygen to oxidize nutrients and to obtain energy
through respiration generates superoxides, hydrogen peroxide, and the highly reactive hydroxyl radicals18. Thus,
microorganisms encounter potentially lethal levels of these ROS due to oxidative burst of phagosomes as well as
from their normal aerobic metabolism27. In fact, NADH primarily serves for the generation of ATP, however, its
oxidation during aerobic respiration is responsible for the generation of most of the endogenous ROS28. Increased
metabolite flow through the glyoxylate shunt instead of the TCA cycle reduces the amount of NADH production
from glucose. The reduction in NADH generation decreases total ROS as superoxides generated from cellular
metabolism will be minimal18. Further, in contrast to eukaryotes, isocitrate dehydrogenase (IDH) in bacteria that

Figure 4. Quantitation of bacterial burden in the spleen upon oral infection of birds with WT or Δms strains of
S. Typhimurium: the spleen was homogenized in sterile PBS and 100 μl homogenate was plated on HEA plates.
The CFU per spleen was calculated after 7 and 14 days post infection (dpi). The data are presented as mean ± SE
(n = 5). *p < 0.05.

Figure 5. Analysis of bacterial burden in the liver upon oral infection of birds with WT or Δms strains of S.
Typhimurium: the liver (100 mg) was homogenized in sterile PBS and 100 μl homogenate was plated on HEA
plates. The CFU per gram of liver was calculated after 7 and 14 days post infection (dpi). The data are presented
as mean ± SE (n = 5). *p < 0.05.
are capable to grow in acetate is linked to NADP rather than NAD. During oxidative stress, NADH functions as the major nicotinamide nucleotide reductant. Due to greater reactivity with Fe³⁺ generated by Fenton’s reaction, the NADH levels rapidly get depleted in comparison to NADPH. In agreement with the above hypothesis parquat stressed E. coli showed increased production of acetate and flux in the glyoxylate cycle consequently increased TCA cycle intermediates following phagocytosis have been observed in fungal pathogens like Paracoccidioides brasiliensis, Penicillium marneffei, Aspergillus fumigatus, Cryptococcus neoformans and Candida albicans as well as in several bacterial pathogens like P. aeruginosa, M. tuberculosis (Munoz-Elias and McKinney 2005) and R. equi. The microorganisms entrapped within the precarious niche of the phagolysosomes require to utilize available carbon sources for their survival under nutrient limiting and other stress conditions. Phagocytes are rich in fatty acids which might induce the activation of the enzymes of the glyoxylate cycle in the phagosome. A study suggested the import of fatty acids derived from host triacylglycerol derldehydes (TAGs) in M. tuberculosis. Further, in S. Typhimurium, the genes for lipid metabolism and glyoxylate cycle are shown to be essential for colonization in mouse tissues like caecum, Peyer’s patches, mesenteric lymph nodes, spleen, and liver. In a co-culture experiment with macrophages, the genes for lipid import, β-oxidation, and glyoxylate cycle were found to be necessary for the survival of S. Typhimurium. Thus, an intriguing possibility of the generation of simple compounds within phagocytes may be furnished by the breakdown of fatty acids via β-oxidation, which results in acetyl-CoA, a substrate for glyoxylate cycle.

In chicken, extensive bacterial multiplication takes place in the caecum. After that, the bacteria reach the spleen, liver, and bone marrow. Earlier, we have demonstrated that malate synthase contributes to the colonization of S. Typhimurium in poultry caecum. Spleen and liver are the important organs involved in immune responses against S. Typhimurium. It has been suggested that replication of S. Typhimurium in the SCV of a single phagocyte in either the liver or spleen is limited during acute infection, however, the bacterial numbers in the liver increase by spreading from one to nearby other phagocytes, thereby forming a focus of infection. Thus, although S. Typhimurium grows poorly in infected phagocytes, it grows in high numbers in the numerous foci of the liver. These observations suggest that the metabolic substrate availability in the phagosome may evolve over the course of infection, with an increasing dependence on fatty acid and acetate utilization occurring during chronic infection. S. Typhimurium modulates splenic functions in ways such that the spleen serves as a safe haven for this bacterium. Next, we assessed the contribution of malate synthase in the survival of S. Typhimurium in poultry spleen and liver. The dissemination of Δms strain to the spleen (p < 0.001) and liver (p < 0.01) was lower than that of the WT strain of S. Typhimurium (Figs. 4 and 5). The involvement of icl as a virulence determinant has been observed in various pathogenic organisms like M. tuberculosis (Munoz-Elias and McKinney 2005), R. equi and P. aeruginosa. In S. Typhimurium, icl is required for chronic infection but not for acute lethal infection in mice.

To survive within the host, bacterial pathogens including S. Typhimurium must have a series of stress management response systems. The existence of several metabolic pathways provides metabolic flexibility which helps in the utilization of various metabolites/substrates available at a particular niche. It has been shown that the complete tricarboxylic acid (TCA) cycle operates during infection of mice with S. Typhimurium. The glycolytic pathway is required for intracellular replication of S. Typhimurium in mice and macrophages and that glucose is the major sugar utilized by S. Typhimurium during infection of macrophages. Thus, to replenish glucose and other TCA cycle intermediates during nutrient limiting and stress conditions, the glyoxylate cycle plays a pivotal role to provide the necessary energy required to replicate and survive within the host.

These observations suggest that metabolic substrate availability in the host may evolve over the course of infection, with an increasing dependence on fatty acid and acetate utilization occurring during chronic infection. Specifically, these enzymes could add potential metabolic flexibility to redox metabolism by effectively decoupling catabolic carbon flow from NADPH formation that would occur in parallel with tricarboxylic acid cycle. Therefore, in the case of S. Typhimurium, the central enzymes of the glyoxylate cycle might be required during different stages to accomplish its metabolic and energy requirements and aid in its survival under oxidative stress conditions.
Experimental procedures

Ethical statement. All animal experiments were approved by the Institutional Animal Ethics Committee (IAEC), Indian Council of Agricultural Research-Indian Veterinary Research Institute (ICAR-IVRI), Izatnagar, India with the approval file No. F.1.53/2012-13-J.D.(Res). All animal experimentations were performed in accordance with the guidelines and regulations of IAEC, ICAR-IVRI, Izatnagar, India.

Bacterial strains and culture media. *Salmonella enterica* subspecies *enterica* serovar Typhimurium strain 5591 (*S. Typhimurium*), a poultry isolate, was procured from the repository of National *Salmonella* Centre-Veterinary, Indian Veterinary Research Institute (IVRI), Izatnagar, India. *ms* gene deletion mutant strain in *S. Typhimurium* (Δ*ms* strain) was cured as described earlier and confirmed by PCR. Bacteriological media like Luria Bertani (LB) agar, LB broth, and Hektoen Enteric agar (HEA) were procured from HiMedia Laboratories Pvt. Ltd., Mumbai, India. Hank’s Balanced Salt Solution (HBSS) was prepared as per standard protocol.

In vitro growth analysis of *S. Typhimurium* and Δ*ms* strains in LB broth and M9 media. Isolated colonies of *S. Typhimurium* and Δ*ms* strains were inoculated in 10 ml of LB broth and grown at 37 °C with shaking at 180 rpm. Overnight grown cultures were diluted (@ 1:100) in fresh 50 ml of LB media and grown on a shaker incubator. Aliquots were withdrawn at every one hour of interval and the optical densities were measured at 600 nm.

The growth of *S. Typhimurium* and Δ*ms* strains were also assessed in M9 minimal media supplemented with acetate or glucose as sole carbon source. Briefly, overnight LB grown cultures were pelleted and washed with M9 minimal media. The washed pellets were then grown in M9 media supplemented with 0.4% acetate or 0.4%glucose.

Evaluation of susceptibilities of *S. Typhimurium* and Δ*ms* strains to HOCl. Overnight grown cultures of *S. Typhimurium* and Δ*ms* strains were exposed to various concentrations of HOCl (sodium hypochlorite, NaOCl, Sigma). Following 2 h of incubation at 37 °C/180 rpm, the cultures were serially diluted and plated on HEA plates. Colonies were enumerated following overnight incubation of the plates.

Evaluation of the susceptibilities of *S. Typhimurium* and Δ*ms* strains to neutrophils. Neutrophils were isolated by a double density centrifugation method using Histopaque 1119/1077 (Sigma-Aldrich, USA). The cells were washed twice with phosphate-buffered saline and once with HBSS devoid of Ca2+/Mg2+ salts (HBSS –) at 250×g for 10 min. The total numbers of viable cells were enumerated by Trypan blue dye exclusion method and adjusted to a concentration of 2 × 10⁶ cells/ml using HBSS (–).

*Salmonella* strains were pelleted, washed, and suspended in HBSS. The neutrophil and bacterial suspensions were mixed at a multiplicity of infection (MOI) of 1:10 (cell:bacteria). The mixture was incubated for 15 min at 37 °C and 5% CO2 in a humidified chamber. To determine the actual MOI, the bacterial suspensions were serially diluted and plated on agar media. Following 15 min of neutrophil-bacterial incubation, the suspensions were centrifuged at 13,000 rpm for 3 min. The supernatant was discarded and the pellet was lysed by 0.1% Triton X-100 for 5 min. Lysates were serially diluted and plated on HEA plates. Colonies were enumerated following overnight incubation of the plates.

Expression analysis of *ms* in *S. Typhimurium*. The expression analysis of the *ms* gene was carried out by qRT-PCR. Neutrophils were infected with *S. Typhimurium* for 15 min as described in the above section. RNA from the harvested pellet was isolated by Trizol reagent. RNA isolated from LB broth grown cultures was served as a control. qRT-PCR was performed according to the protocol as described in Maxima H Minus First Strand cDNA Synthesis Kit (Thermo Scientific). In brief, for a 20 μl reaction, 500 ng of RNA was mixed with 1 μl of random primer (100 pmol), 1 μl of dNTP mix (10 mM), 4 μl of 5× RT buffer, and 1 μl of Maxima H Minus enzyme mix. cDNA was synthesized by incubation of the above mix at 25 °C for 10 min followed by 50 °C for 15 min and termination at 85 °C for 5 min. The expression levels of *ms* gene in each sample were assessed and normalized using DNA gyrase B (*gyrB*) as an internal control. For a 20 μl of qRT-PCR reaction, 0.5 μM of each primer, 2 μl of cDNA, and 10 μl of SYBR Green (Thermo Scientific) were used. The relative fold change in gene expression was determined using the 2−ΔΔCT method. The initial denaturation was carried out at 95 °C for 5 min, followed by 40 cycles consisting of denaturation at 95 °C for 10 s, annealing at 62 °C for 30 s, and data acquisition at 74 °C for 30 s. The amplified products were analyzed on 1.5% agarose gel for assessment of nonspecific amplification (if any) and primer dimers.

Assessment of the contribution of malate synthase in the colonization of *S. Typhimurium* in poultry spleen and liver. All experiments were carried out according to the guidelines of the Institute’s Animal Ethics Committee, ICAR-IVRI, Izatnagar, India and in accordance with the ARRIVE guidelines. The bacterial burdens in the liver and spleen were assessed as described earlier. Briefly, one-day-old chicks were procured from ICAR-Central Avian Research Institute (CARI), Izatnagar, India, and provided with ad libitum feed and water. The birds were screened for the presence of *Salmonella* spp. The *Salmonella* free birds were divided into two groups and orally infected with *S. Typhimurium* or Δ*ms* strain. Following 7, 14, and 21 days post-infection, 5 birds were sacrificed from each group. 100 mg of the liver and whole spleen were homogenized.
in 1 ml of PBS. 100 μl of homogenized samples were plated on HEA plates. The plates were incubated overnight at 37 °C.

**Oxyblot analysis.** Oxidation status of the malate synthase was assessed by OxyBlot™ Protein Oxidation Detection Kit (EMD Millipore). Malate synthase was purified and incubated with 100 mM H₂O₂ as described earlier⁵. Excess H₂O₂ was removed by dialysis. Varying concentrations (2, 1, 0.5 μg) of the H₂O₂ exposed malate synthase samples were denatured with 6% SDS (final) and derivatized with 2, 4-dinitrophenylhydrazine (DNPH). The derivatized samples were then resolved on 10% SDS gel and electrophorrted onto the nitrocellulose membrane. Following blocking the blot was incubated in anti-DNPH antibodies and developed as mentioned in a protocol described elsewhere⁵.

**Western blotting analysis.** S. Typhimurium was cultured in M9 minimal media supplemented either with glucose or acetate. The late stationary phase grown cultures were pelleted and washed with ice cold PBS. The pellets were then lysed by BugBuster Master Mix (EMD Millipore Corp, USA) and unbroken cells were removed by centrifugation. Total proteins in the clarified supernatants were estimated by the Pierce BCA Protein Assay Kit (Thermo Scientific, USA). Fifteen micrograms of cell free lysates were resolved on SDS-gel and transferred to PVDF membranes. Following blocking, the membranes were incubated with anti-MS hyper immune sera (1: 50,000 dilutions in PBS-Tween 20). Following washing, the membrane was washed in anti-rabbit IgG conjugated with alkaline phosphatase (Sigma, at a dilution of 1: 15,000 in PBS-Tween 20) and developed by using NBT and BCIP as substrate⁶.

**Statistical representation.** The graphical representation of the data was done by using Microsoft Excel software and the analysis of results was done by one way analysis of variance (ANOVA).

**Data availability**

All data generated or analysed during this study are included in this published article [and its supplementary information files].

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Author contributions
R.S., S.A., H.B.S. and S.P. performed the experiments. R.S., S.K.B. and M.M. performed the analysis and wrote the paper. All authors reviewed the manuscript.

Competing interests
The authors declare no competing interests.

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