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Paraatha Durgai Amman 108, Lord Durgai in 108 Potri, What is Durgai Amman 108 Antsathas in English - The Hindu Paraatha Durgai Amman 108, Lord Durgai in 108 Potri, What is Durgai Amman 108 Antsathas in English - The Hindu; ? The Indian Express is now on Telegram. Click here to join our channel (@indianexpress) and stay updated with the latest headlines For all the latest India News, download Indian Express App. due to small volumes of measurement material and the need to use a relatively large volume of buffer solution. This however led to a high level of fibrillation during the DEAE-cellulose chromatography. It is important to note that during this step the proteins were diluted 5-6 times in a buffer solution (the main disadvantage of such a buffer). The use of the concentrated protein solution prepared by filtration through a 3 kDa MWCO membrane (3--5 kDa MWCO) and a smaller volume of the buffer solution to overcome the latter problem would result in a very efficient purification process. These steps might lead to even lower fibrillation level. As a consequence of the purification process and having an apparent molecular weight >50 kDa, the proteins were expected to have a completely different conformation than the native ones $[[@B20]]$. The presence of the conformation disorder-induced β -structures in the protein fraction may explain the observed decrease of the protein refolding capability when the proteins were incubated under denaturing conditions. On the other hand, one can speculate that native conformation during the in vitro reconstitution might be important for the quality and efficiency of protein refolding. The importance of the conformation of the native protein has been previously demonstrated for recombinant bacteriophage T4 lysozyme $[[@B21],[@B22]]$. It has been shown that for protein refolding the presence of the native conformation of the target protein is required and such a conformation has higher solubility than the denatured one. Our observations suggest that CGRP has a different conformation during the in vitro refolding process. In fact, the effective refolding capacity of CGRP was found to be much higher than that of CGRP-GFP. The amounts of correctly folded protein are critical

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