

# Theory of Monomer-Dimer Systems<sup>★</sup>

OLE J. HEILMANN<sup>★★</sup> and ELLIOTT H. LIEB

Department of Mathematics  
Massachusetts Institute of Technology, Cambridge, Massachusetts, USA

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**Abstract.** We investigate the general monomer-dimer partition function,  $P(x)$ , which is a polynomial in the monomer activity,  $x$ , with coefficients depending on the dimer activities. Our main result is that  $P(x)$  has its zeros on the imaginary axis when the dimer activities are nonnegative. Therefore, no monomer-dimer system can have a phase transition as a function of monomer density except, possibly, when the monomer density is minimal (i.e.  $x = 0$ ). Elaborating on this theme we prove the existence and analyticity of correlation functions (away from  $x = 0$ ) in the thermodynamic limit. Among other things we obtain bounds on the compressibility and derive a new variable in which to make an expansion of the free energy that converges down to the minimal monomer density. We also relate the monomer-dimer problem to the Heisenberg and Ising models of a magnet and derive Christoffel-Darboux formulas for the monomer-dimer and Ising model partition functions. This casts the Ising model in a new light and provides an alternative proof of the Lee-Yang circle theorem. We also derive joint complex analyticity domains in the monomer and dimer activities. Our considerations are independent of geometry and hence are valid for any dimensionality.

## I. Introduction

A monomer-dimer system is specified by a graph,  $G$  (also called a lattice in the physics literature), together with a family of weights (or Boltzmann factors) assigned to the edges of  $G$ . The precise definition of a weighted graph is given in Section II, but for the present we shall assume the reader is familiar with the concept. Dimers can be placed on the edges of  $G$  so that no vertex has more than one dimer. Uncovered vertices are called monomers and have a fugacity which we call  $x$ . One can also define related problems, such as the monomer-trimer problem, and although the history of these various problems are intertwined we shall consider only the monomer-dimer problem in this paper.

We shall answer the question whether, as the monomer concentration is varied, a phase transition can occur for an infinite system. Our answer,

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<sup>★★</sup> Permanent address: Kemisk Laboratorium III, the H.C. Ørsted Institute, University of Copenhagen, 2100 Copenhagen, Denmark.