

Entanglement in composite free-fermion systems

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We consider fermionic chains where the two halves are either metals with different bandwidths or a metal and an insulator. Both are coupled together by a special bond. We study the ground-state entanglement entropy between the two pieces, its dependence on the parameters and its asymptotic form. We also discuss the features of the entanglement Hamiltonians in both subsystems and the evolution of the entanglement entropy after joining the two parts of the system.

I. INTRODUCTION

The entanglement properties of free-fermion systems have been the topic of many studies and various different cases have been investigated, see e.g. [1]. In one dimension, these comprise homogeneous critical chains where the ground-state entanglement entropy S varies logarithmically with the size L of the subsystem, and non-critical ones where it approaches a constant for large L . Single defects, both at interfaces [2–10] and in the interior [11–13] were studied, as well as inhomogeneous systems with random [14–17], aperiodic [18] and exponentially decaying [19, 20] couplings or random site energies [21–23]. Finally, systems in external potentials which produce a varying density and surface regions have been investigated [8, 24–30].

In the present work, we look at yet another situation, namely at systems formed from two pieces with different properties. Specifically, we study chains composed either of two critical parts, or a critical and a non-critical one. Such systems have been considered previously in the context of conformal invariance [31–34]. In our case, they are realized in the form of undimerized or dimerized tight-binding models coupled by a special bond. Physically, this corresponds to either two metals with different bandwidths, or a metal and an insulator, and we will use this terminology in the following. In both cases one has two types of single-particle eigenfunctions: those essentially confined to one of the subsystems, and other ones extending through the whole system but having different wavelengths in the two parts. This is the same situation as for a potential step in quantum

mechanics. The occupied single-particle eigenfunctions determine the entanglement, and we study it between the two different pieces of the composite system in its ground state.

For the metal-metal system at half filling, the asymptotic result is very simple. It turns out that only the interface bond matters and one comes back to the defect problem solved previously in [4]. Thus S varies logarithmically and the coefficient c_{eff} is determined by the transmission through the interface. The subleading terms, on the other hand, depend on the asymmetry but one can take this into account by a rescaling of the length and find links to conformal formulae based on the nature of the extended states. Away from half filling, the entanglement is small as long as only the localized states are occupied. It increases, when the extended states come into play, but there are strong variations with the filling which depend on the size and are also connected with the ratio of the bandwidths, which brings an additional length scale into the problem.

For the metal-insulator system at half filling, the entanglement lies systematically between that of the two pure systems. At large sizes, the insulator with its band gap dominates and limits the increase of S . For sizes smaller than the correlation length, however, there are no states in the gap and one observes a logarithmic increase of the entropy. The system is also a good case to compare the entanglement Hamiltonians on both sides. They have the same spectra, but their single-particle eigenfunctions and their explicit forms as hopping models are quite different. Basically, one finds that the features are similar to those of the pure systems on the corresponding side of the chain. For the eigenfunctions this means a decay from the interface into the interior which is slow in the metal and exponential in the insulator [1, 35].

For the metal-metal system we also study the behaviour of the entanglement entropy after a local quench in which the two pieces in their ground states are put together. Here the two Fermi velocities can be seen directly in the time structure and the result can be interpreted in the well-known picture of two emitted particles [36].

The paper is organized as follows. In section 2 we describe the set-up and give the basic formulae. In section 3 we investigate the metal-metal case by forming and diagonalizing numerically the correlation matrix. We show the entanglement entropy and discuss its scaling behaviour and filling dependence together with some entanglement spectra. In section 4 we consider the metal-insulator case at half filling and compare all relevant quantities with those of the pure systems. In section 5 we present the time evolution of S after connecting two metallic systems and Section 6 contains a summary. Finally, some analytical results for the metal-metal system are given in an appendix.

II. SETTING AND BASIC FORMULAE

We consider open chains of free fermions with nearest-neighbour hopping and $2L$ sites. The hopping is different in the left and right half and also between both parts. The Hamiltonian is

$$H = -\frac{1}{2} \sum_{n=-L+1}^{L-1} \hat{t}_n (c_n^\dagger c_{n+1} + c_{n+1}^\dagger c_n) \quad (1)$$

For the metal-metal system

$$\hat{t}_n = \begin{cases} t_1 & : n < 0 \\ t_0 & : n = 0 \\ t_2 & : n > 0 \end{cases} \quad (2)$$

and one has three parameters in the problem. However, by a rescaling of H , one can always achieve $t_1 t_2 = 1$. We will assume this in the following and write the quantities as $t_1 = \exp(\Delta)$, $t_2 = \exp(-\Delta)$ and $t_0 = \exp(\Delta_0)$. Moreover, we will always consider positive Δ , i.e. $t_1 > t_2$. In some places we also use the ratio $r = t_2/t_1$.

For vanishing coupling ($t_0 = 0$), the two parts of the chain have single-particle energies

$$\omega_\alpha = -t_\alpha \cos(q_\alpha), \quad \alpha = 1, 2 \quad (3)$$

with momenta $q_\alpha = \pi m/(L+1)$, $m = 1, 2, \dots, L$. The corresponding band structure with bands between $\pm t_\alpha$ is shown on the left of Fig. 1. In the coupled system, one has extended states for $|\omega| < t_2$, while outside this region they are confined essentially to the left half-chain. Moreover, one can have two states localized at the interface if t_0 is large enough.

For the metal-insulator system we take

$$\hat{t}_n = \begin{cases} 1 & : n < 0 \\ t_0 & : n = 0 \\ 1 + (-1)^{n+1} \delta & : n > 0 \end{cases} \quad (4)$$

In the insulator, one thus has alternating hopping, two sites per unit cell and the single-particle energies

$$\omega_2 = \pm \sqrt{\cos^2(q_2) + \delta^2 \sin^2(q_2)} \quad (5)$$

where the momenta have to be determined from the boundary condition. Thus there is a gap between $\pm\delta$. The resulting band structure is shown on the right of Fig. 1. For an open chain, there are also states localized at the boundary with exponentially small ω_2 , if the outermost bond

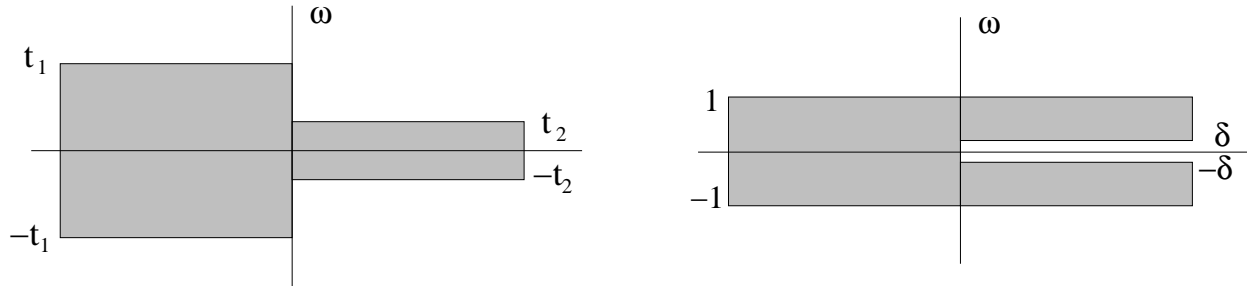


FIG. 1: Schematic band structure of the uncoupled composite systems. Left: Metal-Metal, Right: Metal-Insulator.

is a weak one. The coupled system has extended states for $|\omega| > \delta$ and others, confined essentially to the metal, inside the gap. In this sense, it is still critical.

It is simple to set up the eigenvalue problem for the full systems, and some details are given in the Appendix. However, the matching conditions in the center are difficult to handle in general. Therefore we determine the single-particle eigenvalues ω_q and the corresponding (real) eigenfunctions $\Phi_q(n)$ numerically. The correlation matrix $C_{mn} = \langle c_m^\dagger c_n \rangle$ is then

$$C_{mn} = \sum_{q \in F} \Phi_q(m) \Phi_q(n) \quad (6)$$

where the sum extends over the states q in the Fermi sea. Restricting C_{mn} to the chosen subsystem (either the left or the right half-chain), its eigenvalues ζ_k give the single-particle eigenvalues $\varepsilon_k = \ln[(1 - \zeta_k)/\zeta_k]$ of the free-fermion entanglement Hamiltonian \mathcal{H} in

$$\rho = \frac{1}{Z} \exp(-\mathcal{H}) \quad (7)$$

where ρ is the reduced density matrix [37]. From them, the entanglement entropy S follows as

$$S = \sum_k \ln(1 + e^{-\varepsilon_k}) + \sum_k \frac{\varepsilon_k}{e^{\varepsilon_k} + 1} \quad (8)$$

In addition to the eigenvalues ζ_k resp. ε_k , we also determine the corresponding eigenfunctions φ_k and construct \mathcal{H} in section 4.

III. METAL-METAL SYSTEM

We first consider half-filled systems where the Fermi level is in the middle of the bands. The resulting entanglement entropies for $t_0 = 1$ and three values of the parameter Δ are shown in Fig. 2, both for even and for odd L . One sees that the typical increase with the size, known for the

homogeneous case $\Delta = 0$, persists in the composite systems. The plot against $\ln L$ shows that also the asymptotic law $1/6 \ln L + k$ is unchanged. However, the value of k becomes smaller and the finite-size effects, in particular the even-odd alternation, increase dramatically as Δ becomes larger. For $\Delta = 3$ (not shown), the values of S are initially close to zero and to $\ln 2$, respectively. In this case, the ratio of the bandwidths $t_2/t_1 \simeq 1/400$ is already very small, the states localized on the left drop rapidly on the right, while the extended states have very small amplitudes on the left (see the Appendix). For even L , this leads to a small entanglement which only increases as more states come into play at larger L . The value $\ln 2$ for odd L is a consequence of the particle-hole symmetry of the problem which, in this case, forces one of the ε_k to vanish.

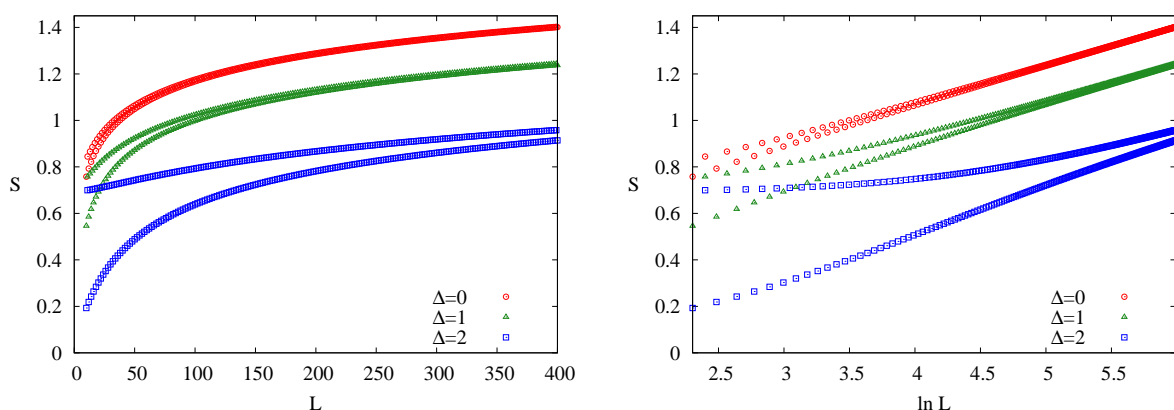


FIG. 2: Entanglement entropy S for $\Delta_0 = 0$ and three values of Δ . Left: as function of L . Right: as function of $\ln L$. The upper (lower) curves correspond to odd (even) values of L .

The function $S(L, \Delta)$ has an intriguing universal behaviour. If one defines a length $L^* = L/\alpha$ and chooses the scale factor α properly, one can achieve $S(L^*, \Delta) = S(L, 0)$, i.e. all curves collapse on the one for the homogeneous system. Written differently, one has

$$S(L, \Delta) = S(\alpha L, 0) \quad (9)$$

In the asymptotic region, where one has straight lines in a logarithmic plot, this feature is not surprising: A rescaling can always generate the necessary shift to make the curves coincide. However, the phenomenon is not restricted to this region. As shown in Fig. 3 on the left, it also holds for small sizes where there is still curvature in the graph. The variation of α with Δ is given on the right hand side of Fig. 3. One sees that the results determined from different (even) L agree very well and that α becomes rapidly smaller as Δ increases. The curve can be described approximately by $1/\text{ch}^2 \Delta$. Note that the relation (9) can only be applied to values of L such that αL is larger than 1.

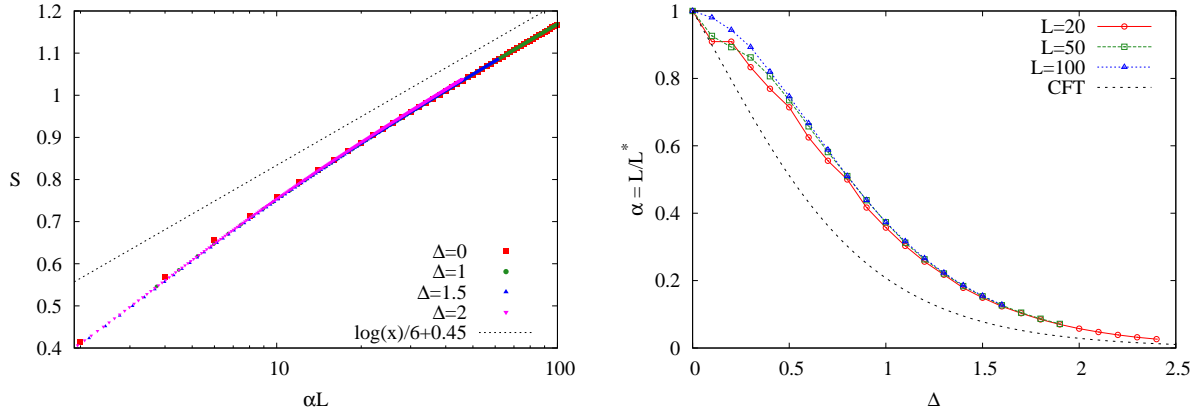


FIG. 3: Scaling behaviour of the entanglement entropy. Left: S as a function of the effective length αL for four values of Δ (logarithmic scale). Right: Variation of $L/L^* = \alpha$ with Δ . The dotted line is the conformal factor in (10).

At this point, an observation from the appendix is helpful. Namely, near the Fermi level the wavefunctions in the right subsystem are the same as for a homogeneous system of total length $L(1+r)$. Thus, ignoring the states outside the narrow band, the correlation matrix on the right is the same as for an unsymmetrically divided homogeneous chain and one can invoke the conformal result for the entropy [38] extended in [39] to include $1/L$ corrections. This gives

$$S_{FC} = \frac{1}{6} \ln z - \frac{(-1)^\ell}{z} + \text{const.}, \quad z = \frac{4}{\pi} L(1+r) \sin \frac{\pi \ell}{L(1+r)}, \quad (10)$$

with $\ell = L$. The r -dependent factor in z provides a similar rescaling of L as α , and a corresponding shift of the curves. It turns out, however, that this is not enough to make them coincide. Rather, one has a residual difference $S - S_{FC} = S_I(\Delta)$, which one can attribute to the interface, and which rises smoothly from 0 to about 0.13 as Δ increases from 0 to 3. It has its origin in the neglected exponentially decaying states below the narrow band.

This can be seen clearly, if one considers a partition of the system with $\ell < L$ sites on the right and calculates the entanglement between this subsystem and the remainder. As demonstrated in Fig. 4, the expression (10) then fits the data very well as soon as one moves away from the interface by a few lattice sites. One can also consider a division located in the left part of the system where the extended wave functions are the same as in a homogeneous system with total length $L(1+1/r)$, but in this case the conformal formula does not work well because the additional states are more important. These results also show that the symmetrical division is actually a somewhat special case.

We now turn to the effect of t_0 . If one varies the central bond in a homogeneous chain ($\Delta = 0$),

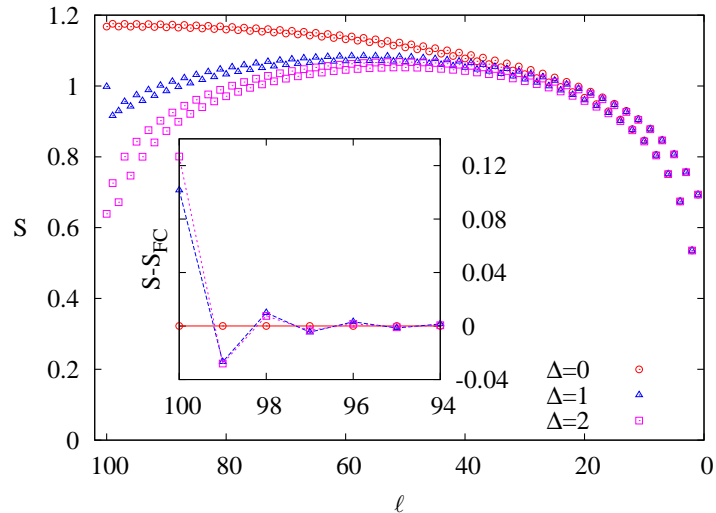


FIG. 4: Entanglement entropy between the outermost ℓ sites of the right subsystem and the remainder for $L = 100$ and three values of Δ . The inset shows the deviation from the formula (10).

the asymptotic behaviour of S is

$$S = \frac{c_{\text{eff}}}{6} \ln L + k \quad (11)$$

where c_{eff} depends on t_0 and is given by an explicit, though lengthy formula in which only the transmission coefficient at the Fermi energy $T \equiv s^2$ enters [4]. But the calculation in the Appendix shows that T is independent of Δ and given by

$$T = \frac{1}{\text{ch}^2 \Delta_0} \quad (12)$$

Therefore one expects the result of the homogeneous problem also in the composite systems. This is, in fact, the case and demonstrated in Fig. 5 on the left, where results obtained by fitting the data between $L = 100$ and $L = 400$ to (11) plus a $1/L$ term are collected. For $\Delta = 2$, the average between even and odd sizes was taken, while for $\Delta = 1$ this was unnecessary. This verifies the Fermi-edge character of c_{eff} also in the present case.

The constant k , on the other hand, depends on Δ , as was found above already for the special case $t_0 = 1$. It is shown in Fig. 5 on the right. The values for odd L are always larger than those for even L with the maximal difference of $\ln 2$ appearing for $t_0 \rightarrow 0$. This is connected with the half-integer particle number in each subsystem for odd L which leads to a kind of singlet state even in the decoupling limit. The minimum in k for even L is also present for $\Delta = 0$ and was also found for a segment in an infinite chain [2].

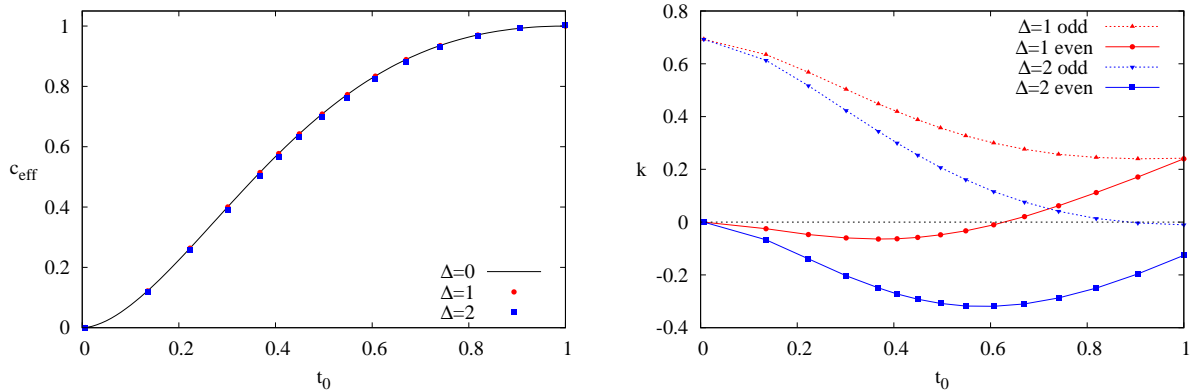


FIG. 5: Effective central charge c_{eff} and constant k in (11) as functions of t_0 for different values of Δ . The line $\Delta = 0$ in the left figure is the theoretical result, see [4].

Finally, we consider the dependence of S on the filling $\nu = N/2L$ of the system, where N is the number of particles. It is shown in Fig. 6 for a fixed length $L = 200$ and four values of Δ . The smooth curve (red) is the result for the homogeneous system and given, up to the small even-odd oscillations [39–42] and a constant, by $1/6 \ln(\sin q_F)$, where $q_F = \pi\nu$ [43]. Turning on Δ , regions of very small entanglement develop for small and large filling, where only states localized on the left are occupied or remain empty. At the same time, the oscillations of S in the central region become slower and slower.

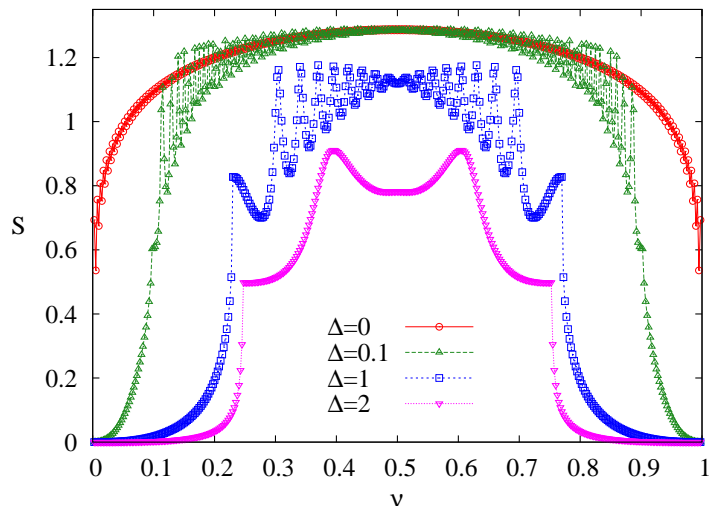


FIG. 6: Entanglement entropy S as a function of the filling $\nu = N/2L$ for $L = 200$ and different Δ .

The origin of these oscillations, at the level of the eigenvalues ε_k , can be seen in Fig. 7. There, the low-lying ε_k are shown for all fillings between 0.2 and 0.8. For the case $\Delta = 2$, shown on the

right, the variation of the ε_k with ν is rather slow and the curves cross zero only at two fillings. At these points, the crossing eigenvalue gives the maximal contribution to S , namely $\ln 2$, and they coincide with the locations of the maxima in Fig. 6. For $\Delta = 1$, shown on the left, the variation is much faster and one has altogether 16 crossings. This is again the number of maxima of S in the center of Fig. 6.

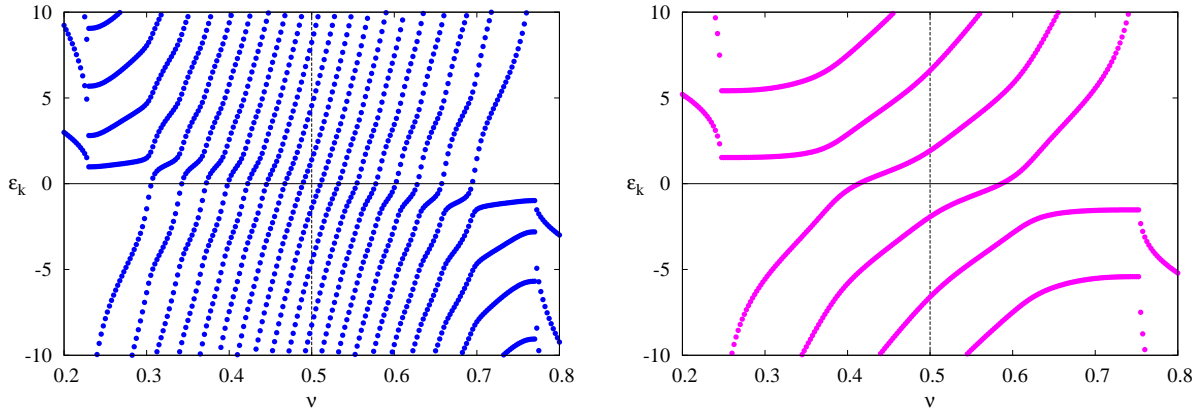


FIG. 7: Variation of the low-lying single-particle eigenvalues ε_k with the filling $\nu = N/2L$ for $L = 200$. Left: $\Delta = 1$. Right: $\Delta = 2$. The colours are the same as in Fig. 6.

These structures can again be understood from the relation to a smaller, unsymmetrically divided homogeneous system. In such a case, if L' is the size of the smaller subsystem, one has L' crossings of the eigenvalues and the effect on S is given by a factor $\sin(q_F(2L' + 1))/\sin q_F$ replacing the alternating sign of the $1/z$ term in (10). This leads to slow oscillations, although less pronounced than the observed ones. However, $L' = rL$ gives $L' = 3.6$ for $\Delta = 2$ and $L' = 27$ for $\Delta = 1$ and thus not the correct numbers (2 and 16), even if one takes the nearest integer. But one can argue that the length rL only refers to the states in the center of the band, while near the edges $\sin k_2$ appears in (26) which is smaller than k_2 by a factor of $2/\pi$. Working with $\tilde{L}' = 2rL/\pi$ gives values 2.3 and 17.2, respectively, which are quite close to the numerical findings. One also arrives at \tilde{L}' by counting the number of levels which the left subsystem contributes to the total number of band states. To accommodate them, one needs just this number of additional sites.

One should mention that one can see basically the same structures also in the energy ω as tiny deviations from the otherwise smooth level spacing of the extended states. In this case, the interpretation is that \tilde{L}' also gives the number of periods of the slowly varying tangent in (26) as k_2 sweeps through the band, and thus the number of possible peculiarities in the allowed momenta.

IV. METAL-INSULATOR SYSTEM

In the following, we always consider half-filled systems. To avoid boundary states in the dimerized subsystem, we work with even L and choose strong bonds $1 + \delta$ at the borders. The entanglement entropies for two relatively small dimerizations and $t_0 = 1$ are shown in Fig. 8, together with those for the pure systems. One sees that the mixed system always has an intermediate entanglement, which is very plausible. Replacing one half of the metal by an insulator reduces the metallic result, while replacing one half of the insulator by a metal increases the insulator result. For $\delta = 0.001$, shown on the left, all three curves look similar and there is no sign of a saturation, while for $\delta = 0.01$, shown on the right, all three curves show a clear saturation.

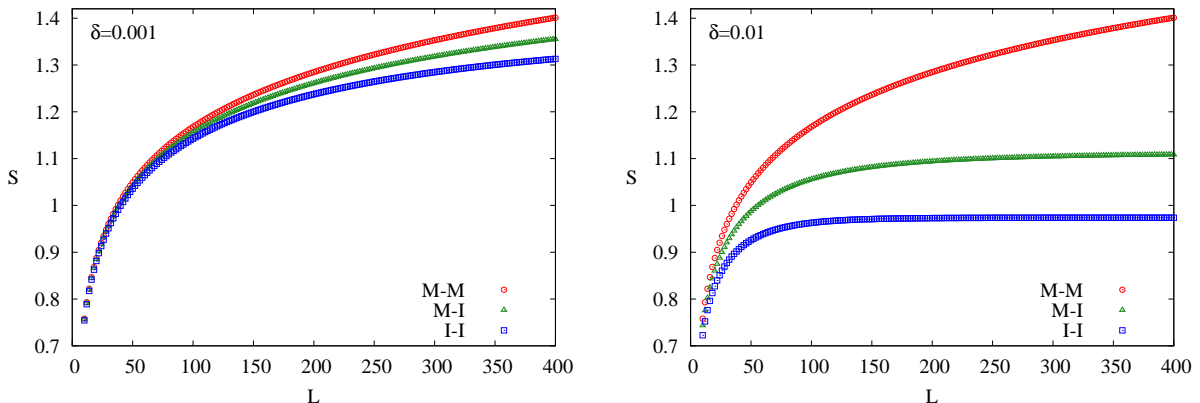


FIG. 8: Entanglement entropies for metal-metal, metal-insulator and insulator-insulator systems as a function of L for two values of the dimerization parameter δ .

while for $\delta = 0.01$ a saturation is clearly visible for the lower ones. This can be interpreted in terms of the correlation length $\xi = 1/\delta$ in the insulator. It is $\xi = 1000$ in the first case so that the non-criticality does not fully show up. In the second case, where $\xi = 100$, it does and the crossover takes place roughly around this value.

In the region $L \ll \xi$, one can fit the curves for the composite system to a form $a \ln L + b$ and finds values of a quite close to $1/6$. This can also be done for $t_0 < 1$ and gives curves for c_{eff} which lie somewhat below the metallic one shown in Fig. 5, but approach it as δ goes to zero. In the opposite limit, $L \gg \xi$, the entropy saturates and for $t_0 = 1$ one finds the asymptotic law $S \simeq 1/6 \ln \xi + \text{const.}$ as for the pure insulator but with a larger constant. To obtain the coefficient $1/6$ accurately from not too large ξ , one has to include subleading corrections of the form $(d \ln \xi + e)/\xi$ which one knows in the pure case from the connection to a transverse Ising chain sketched below. Again, the analysis can be extended to $t_0 < 1$, and in this case one finds c_{eff} to an accuracy of 3-4 digits.

On the level of the wavefunctions entering the correlation matrix, the (pseudo-) critical be-

haviour can be understood in the following way. For $L < \xi$ one has $|\omega| \sim 1/L > \delta$ and there are no states in the metal with energy in the band gap of the insulator and none in the insulator too close to the band edge. Thus one only has extended wavefunctions which are similar to those in the pure metal. This can be seen analytically or from the numerics.

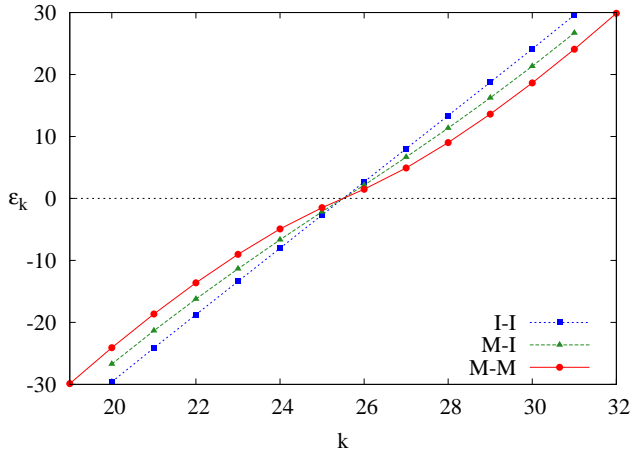


FIG. 9: Eigenvalues ε_k for metal-metal, metal-insulator and insulator-insulator systems with $L = 50$ and $\delta = 0.1$ for the insulator.

It is also instructive to look at the ε_k spectra. In Fig. 9 an example is shown for the case $\delta = 0.1$ and $L = 50$. The lowest curve with the slight bend is the well-known result for the metal, while the highest one is for the insulator and strictly linear as for the transverse Ising chain or the XY model [1]. In fact, one can find a relation between the ε_k for the dimerized hopping model and the transverse Ising chain using the method in [44]. The latter model then has field $h = 1$ and coupling $\lambda = (1 - \delta)/(1 + \delta)$ and the ε_k are

$$\varepsilon_k = (2k + 1)\varepsilon, \quad k = 0, \pm 1, \pm 2, \dots, \quad \varepsilon = \pi I(\lambda')/I(\lambda) \quad (13)$$

where $I(\lambda)$ denotes the complete elliptic integral of the first kind and $\lambda' = \sqrt{1 - \lambda^2}$, see also [45]. The MI result lies in between, with a small break after the first eigenvalue, which becomes more apparent for larger values of δ , when the curve becomes steeper and more linear.

For the spectrum, it does not matter in which of the two subsystems one considers the correlation matrix. Formally, this follows from the property $C^2 = C$ for the full system [46]. However, the eigenvectors of the two reduced matrices, and therefore the corresponding entanglement Hamiltonians \mathcal{H} , are quite different. This is illustrated in Fig. 10 for the lowest $|\varepsilon_k|$ and $\delta = 0.1$. We have plotted $\varphi_k(n)$ for $+\varepsilon_k$ on the left and for $-\varepsilon_k$ on the right, since these two are directly related. Namely, if one partitions C into four blocks according to the location of the sites and φ^1 is an

eigenvector of C^{11} for part 1 with eigenvalue ζ , then

$$\varphi^2 = C^{21}\varphi^1 \quad (14)$$

is an eigenvector of C^{22} for part 2 with eigenvalue $1 - \zeta$.

One sees the typical features of the pure systems on the two sides, namely a slow power-law decay in the metal (left) and a fast one within a distance $\xi = 10$ in the insulator (right). The pure cases are shown for comparison, and one notes that the differences in the composite system are relatively small, lying more in the amplitudes than in the general behaviour.

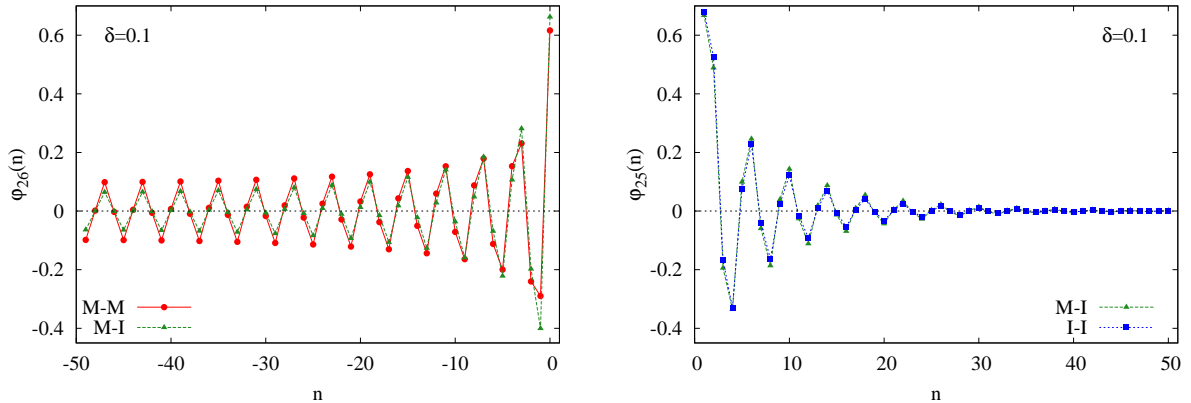


FIG. 10: Eigenvectors of the MI correlation matrix for the smallest $|\varepsilon_k|$ in the metallic (left) and the insulating (right) subsystem for $L = 50$ together with the results for the homogeneous cases (MM and II).

With the eigenvectors, one can explicitly determine the entanglement Hamiltonians \mathcal{H} . They have again the form of hopping models with basically only nearest-neighbour hopping as in H , but the amplitudes \tilde{t}_n increase from the center towards the boundaries.

In the pure metal, this increase is monotonous and the \tilde{t}_n vary as $n(2L - n)$ [1], while in the insulator one has an additional alternation which is coupled to the bond alternation in H . Due to the form of the eigenvectors, the result in the composite system is again close to that of the pure system on the corresponding side. This is shown in Fig. 11 for a system with a total of $2L = 16$ sites, which is the largest size attainable before numerical errors in the large ε_k set in.

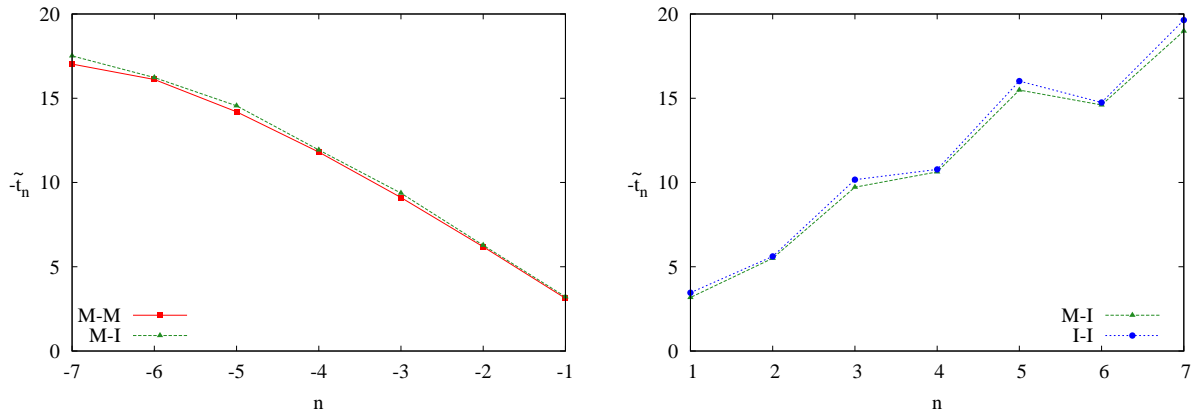


FIG. 11: Nearest-neighbour hopping in the entanglement Hamiltonians for the left and right subsystem as a function of the position together with the results for the pure systems for $L = 8$ and $\delta = 0.1$.

V. EVOLUTION AFTER A QUENCH

In this section, we return to the metal-metal systems and study how the entanglement evolves after one joins the two initially disconnected parts. This type of local quench has been considered repeatedly in the past [11, 47–59]. For the homogeneous case, one finds oscillations of $S(t)$ which one could call “entanglement bursts”, see e.g. [51]. In our case, the situation is more complex, because of the interface and the two Fermi velocities.

The two pieces are coupled at time $t = 0$ by switching on t_0 and the evolution of $C(t)$ is calculated via the Heisenberg operators $c_n(t), c_n^\dagger(t)$. The resulting $S(t)$ is shown in Fig. 12 for two different values of Δ and of t_0 . The system on the left with $\Delta = 0.5$ has roughly the band widths of Fig. 1, while on the right the asymmetry is larger.

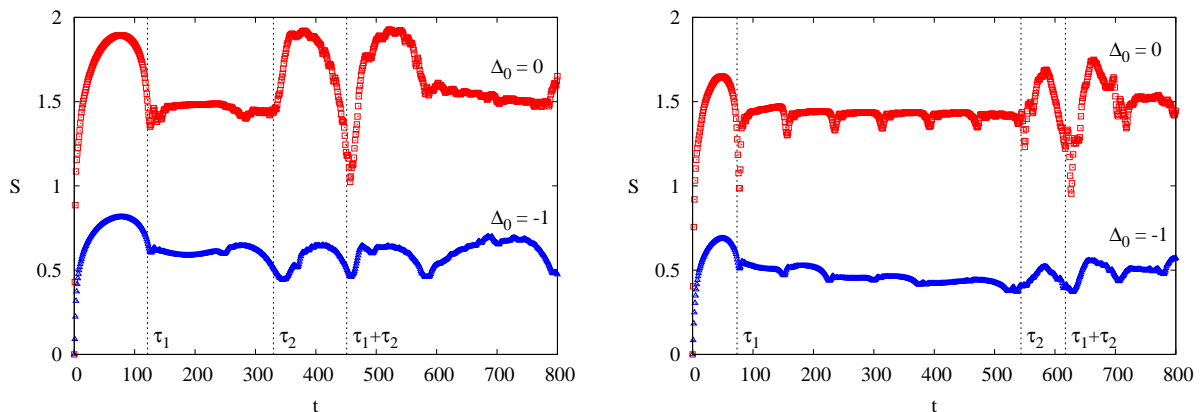


FIG. 12: Evolution of the entanglement entropy S after connecting two metallic systems with $L = 100$. Left: $\Delta = 0.5$. Right: $\Delta = 1$. The times indicated are the theoretical values, see the text.

For $\Delta_0 = 0$ (upper curves) one has the following general features: An initial burst ending with a decrease at a time τ_1 followed by a plateau, another burst at time τ_2 which is a kind of mirror image of the first ending at time $\tau_1 + \tau_2$ and then a rough repetition. For $\Delta = 1$, the plateau is longer and shows six additional structures, whereas for $\Delta = 0.5$ only a single one is visible. The lower curves have roughly the same pattern, but apart from the first maximum the structures are more washed out.

These features can be interpreted in the well-known picture due to Calabrese and Cardy [36], in which a pair of particles is emitted from the junction at time $t = 0$, travelling in opposite directions and spreading the entanglement. In the present case, they have the velocities $v_1 = t_1$ and $v_2 = t_2$ and the corresponding space-time diagram is given in Fig. 13.

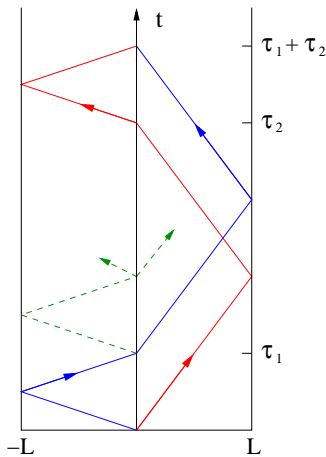


FIG. 13: Trajectories of the particles in the Calabrese-Cardy picture.

After time $\tau_1 = 2L/v_1$, the left particle returns to the center, and if it simply moved into the right subsystem, the entanglement would continue to drop. This happens in a somewhat related situation where the velocities are the same, but the size of the right subsystem is larger [51]. Here, however, there is a probability for reflection at the interface, such that the entanglement rises again and another small “burst” follows. For $\Delta = 1$, where $v_1 = 7.4 v_2$, the left particle can make seven round trips before the right particle returns to the center at time $\tau_2 = 2L/v_2 = \tau_1/r$. These are very clearly visible in the Figure. There are, however, some remarks to make. According to (12) there is no reflection right at the Fermi level for $\Delta_0 = 0$. The effect can therefore only come from somewhat slower particles away from it. In fact, the numerical values for τ_1 and τ_2 are somewhat larger than the theoretical ones, which supports this argument. Also, one would expect the reflection effect to become weaker with each cycle, which is only barely the case. On the other hand, for $\Delta_0 = -1$

one has $R = 0.58$ at the Fermi level and the effect should be stronger, which is also not the case. Nevertheless, the picture gives a good overall description.

One could presume that at least the first burst can be described by the conformal result [51]

$$S_h(vt) = \frac{c}{3} \ln \left| \frac{2L}{\pi} \sin \frac{\pi vt}{2L} \right| + \text{const.} \quad (15)$$

valid for a homogeneous system with particle velocity v . Thus we tried the Ansatz

$$S(t) = [S_h(v_1t) + S_h(v_2t)]/2 + \text{const.} \quad (16)$$

which, on the scale of Fig. 12, fits the data for $\Delta_0 = 0$ very well. However, a closer look at the final decrease shows that it comes too early since, as mentioned, the value of τ_1 is too small. Finally, we note that the first-burst results for a homogeneous but unequally divided system with the same round-trip times $\tau_1 = 2L_1$ and $\tau_2 = 2L_2$ lie above ours, and the difference increases with Δ . Thus the two problems are not trivially connected.

VI. SUMMARY

We have considered the entanglement in fermionic chains composed of two different halves. In solid-state terminology, they were metals and insulators, while in statistical-physics terms they corresponded to critical and non-critical systems. The metal-metal system at half filling showed the same logarithmic behaviour of S as a homogeneous chain with a defect. In the subleading terms, the asymmetry showed up, but the entropy had a remarkable scaling property. Its filling dependence, finally, showed oscillatory behaviour again linked to the asymmetry. Both features could be understood from the nature of the extended eigenfunctions which are the same as in homogeneous systems with different lengths.

The difference in the Fermi velocities showed up directly in the quench experiment where two half-filled metals were joined and the entanglement was monitored. In that case, the value of r determined, how many cycles one sees in $S(t)$, before a certain return to the initial situation takes place. This pattern would occur in unsymmetrically divided homogeneous systems only with additional defects.

The metal-insulator system was somewhat simpler. Its entanglement properties were seen to lie always between the two pure systems and as a function of the size, one has a crossover from an initial critical behaviour with a logarithmic increase of the entanglement to a saturation typical for a non-critical system with finite correlation length. It also provided an instructive example

for entanglement Hamiltonians which are quite different for the two subsystems although they have the same spectra, as they must. One should note that the dimerized hopping model used for the insulator, and investigated already in [45], plays a central role in the theory of polyacetylene [60, 61]. In that sense, we studied a particular polymer system.

We considered the geometry with open ends, because then one has only a single interface. However, one could equally well look at rings. For composite transverse Ising models this was done first in [31, 32] but without particular interface bonds and with a view on the spectra. Obtaining the entanglement entropy is more complicated, and we could only give an analytical result for the asymptotic behaviour of the metal-metal system.

While we focussed on the entanglement entropy, it is known that the particle-number fluctuations in the subsystems behave similarly [62]. Thus, in a homogeneous metal, the prefactor of the $\ln L$ term in the fluctuations is given by $T/2\pi^2$ with the transmission coefficient T [8] and varies qualitatively like c_{eff} . The same result is found in the composite systems, and this could offer a way to access the entanglement experimentally.

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Appendix: Some formulae for the metal-metal system

Let $\bar{\phi}(n)$ and $\phi(n)$ denote the wave function at site n on the left and right, respectively. The bulk solutions on the two sides are waves with momenta q_1 and q_2 and energy $\omega = -t_1 \cos q_1 = -t_2 \cos q_2$. At the interface, the equations are

$$t_0 \bar{\phi}(0) = t_2 \phi(0) \quad , \quad t_1 \bar{\phi}(1) = t_0 \phi(1) \quad (17)$$

For a wave coming in from the left and being partially reflected and transmitted, the amplitudes are

$$\begin{aligned} \bar{\phi}(n) &= A_1 \exp(iq_1 n) + B_1 \exp(-iq_1 n) \\ \phi(n) &= A_2 \exp(iq_2 n) \end{aligned} \quad (18)$$

Inserting this into the relations (17), one finds for the reflection coefficient

$$R = \left| \frac{B_1}{A_1} \right|^2 = \frac{\operatorname{ch} 2\nu - \cos(q_1 - q_2)}{\operatorname{ch} 2\nu - \cos(q_1 + q_2)} \quad (19)$$

where the quantity ν is defined as $\exp(2\nu) = t_0^2/t_1t_2$. In our parametrization, where $t_1t_2 = 1$, one has $\nu = \Delta_0$. The transmission coefficient follows from $T = 1 - R$ and can be written

$$T = \frac{\sin q_1 \sin q_2}{\operatorname{sh}^2 \nu + \sin^2((q_1 + q_2)/2)} \quad (20)$$

If the left and right subsystems are identical, $q_1 = q_2 = q$, and one finds the result for a bond defect in an otherwise homogeneous chain. The same holds in a general system if one is in the middle of the bands. Then $\omega = 0$, $q_1 = q_2 = \pi/2$ and one has the result

$$T = \frac{1}{\operatorname{ch}^2 \nu} \quad (21)$$

Due to the factor $\sin q_2$ in (20), the transmission vanishes at the edges of the narrow band.

The extended eigenfunctions of H have the form

$$\begin{aligned} \bar{\phi}(n) &= A_1 \sin q_1(n + L) \\ \phi(n) &= A_2 \sin q_2(n - L - 1) \end{aligned} \quad (22)$$

and the conditions (17) lead to

$$\frac{\sin q_1(L + 1) \sin q_2(L + 1)}{\sin q_1 L \sin q_2 L} = \frac{t_0^2}{t_1 t_2} \quad (23)$$

which, together with $t_1 \cos q_1 = t_2 \cos q_2$, determines the allowed momenta. Setting $q_\alpha = \pi/2 + k_\alpha$ and assuming L even, it takes the form

$$\frac{\cos k_1(L + 1) \cos k_2(L + 1)}{\sin k_1 L \sin k_2 L} = \frac{t_0^2}{t_1 t_2} \quad (24)$$

The relation between the k_α reads

$$\sin k_1 = r \sin k_2, \quad r = t_2/t_1 \quad (25)$$

and for small r simplifies to $k_1 = r \sin k_2$. Inserting this into (24) and setting $L + 1 \simeq L$ in the numerator then gives

$$\cot(k_2 L) = \frac{t_0^2}{t_1 t_2} \tan(rL \sin k_2) \quad (26)$$

which contains only the momentum k_2 . The solutions can be discussed graphically, but one sees directly that besides L a second length $L' = rL$ appears. If it is large enough, the tangent

completes several cycles as k_2 varies, and this can lead to additional features in the momenta or in the amplitude ratio which is, again with $L + 1 \simeq L$,

$$\frac{A_2}{A_1} = -\frac{t_1}{t_0} \frac{\cos(rL \sin k_2)}{\sin k_2 L} \quad (27)$$

If k_2 is small, $k_1 = rk_2$ for all $r < 1$ and (26) for $t_0^2/t_1 t_2 = 1$ can be written in the two forms

$$\tan(k_2 r L) \tan(k_2 L) = 1, \quad \tan(k_1 L) \tan(k_1 L/r) = 1 \quad (28)$$

Both equations are the same as for a *homogeneous* chain built up from two pieces with different lengths and can therefore be solved explicitly. For k_2 , the homogeneous system has length $L + L'$, while for k_1 it has length $L + L''$, where $L'' = L/r$. Thus the small momenta are equidistant, i.e.

$$k_2 = \frac{\pi}{2(L + L')} (2n + 1), \quad n = 0, \pm 1, \pm 2, \dots \quad (29)$$

Using (28) in (27), one finds $(A_2/A_1)^2 = t_1^2 = 1/r$, i.e. the amplitudes are larger on the right than on the left. Including normalization to leading order, this gives prefactors $A_1^2 = 2/(L + L'')$ and $A_2^2 = 2/(L + L')$ which correspond exactly to these chain lengths.

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