Quantum Periods and Spectra in Dimer Models and Calabi-Yau Geometries

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Abstract

We study a class of quantum integrable systems derived from dimer graphs and also described by local toric Calabi-Yau geometries with higher genus mirror curves, generalizing some previous works on genus one mirror curves. We compute the spectra of the quantum systems both by standard perturbation method and by Bohr-Sommerfeld method with quantum periods as the phase volumes. In this way, we obtain some exact analytic results for the classical and quantum periods of the Calabi-Yau geometries. We also determine the differential operators of the quantum periods and compute the topological string free energy in Nekrasov-Shatashvili (NS) limit. The results agree with calculations from other methods such as the topological vertex.

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1 Introduction and Summary

The developments of various prosperous topics in mathematics and physics often intersect each others. Topological string theory on Calabi-Yau manifolds has been a fruitful branch of superstring theories that encompass many recurring themes in mathematical physics, see e.g. [1]. In the seminar work [2], Nekrasov and Shatashvili (NS) proposed a connection between the partition function of Seiberg-Witten gauge theory on Ω background and certain quantum integrable systems. In the NS limit, we set one of the two Ω deformation parameters to vanish and identify the other as the Planck constant of the quantum system. This relation can be uplifted to five dimensions, where the partition functions are computed by refined topological string theory on corresponding Calabi-Yau spaces. The topological free energy in the NS limit can be viewed as a quantum deformation of the prepotential, and is computed similarly by promoting the periods of the Calabi-Yau geometries to quantum periods [3, 4, 5, 6]. More examples in Seiberg-Witten theories can be found in [7, 8, 9, 10, 11]. The quantization conditions of the quantum system are formulated as the Bohr-Sommerfeld quantization condition where the phase volumes are computed by quantum periods. In the five dimensional case, the quantum systems are often known as relativistic models due to the exponential kinetic and potential terms in the Hamiltonians from
quantizing the mirror curves of the local Calabi-Yau spaces. Inspired by earlier works [12, 13, 14], some novel non-perturbative contributions to the quantization conditions are conjectured in [15, 16]. Various aspects of the quantization conditions, including complex value Planck constant, resurgence, wave functions, etc are further explored in e.g. [17, 18, 19, 20, 21]. The non-perturbative parts of the two types of exact quantization conditions in [15, 16] are related by certain constrains on the BPS invariants known as the blowup equations [22][23]. The blowup equations originally come from studies of Seiberg-Witten gauge theories [24] (see also [25, 26]), but have now become a very effective tool for computing topological string amplitudes on various Calabi-Yau manifolds [27, 28, 29, 30, 31]. The exact quantization conditions have also been applied to related condensed matter systems, e.g. in [32, 33, 34, 35].

Most examples of the early studies focus on geometries with mirror curves of genus one. The quantum periods and quantization conditions for quantum systems corresponding to mirror curves of higher genus were subsequently considered, in e.g. [36, 37, 22, 38, 39]. A particularly interesting class of quantum integrable systems can be constructed by dimer models on torus [40], and the quantization conditions are studied in [41, 42]. The dimer models in this paper also correspond to local toric Calabi-Yau geometries and the mirror curves are encoded in the data of the bipartite dimer graphs. Some of Calabi-Yau spaces geometrically engineer 5d supersymmetric gauge theories, which are uplifts of the 4d $SU(N)$ Seiberg-Witten theories considered in [7]. There are a number of commuting Hamiltonians, and the multiple quantization conditions can be similarly derived from topological string free energy in the NS limit on the corresponding Calabi-Yau spaces. The studies in [41, 42] mostly focus on numerical tests of the non-perturbative quantization conditions. However, in order to have a more insightful understanding of the interconnections between various subjects here, it is better to have some analytical results. In this paper we develop some analytic approaches to the problem, though mostly focusing on the perturbative aspects.

The paper is organized as followings. In Section 2 we review the constructions of dimer models, and derive Hamiltonians of the quantum integrable systems based on previous literatures. We shall study some examples with genus two mirror curves and correspondingly two commuting dynamical Hamiltonians. In Section 3 we study the perturbative quantum spectra of the Hamiltonians around minimal points of the phase space. A useful technical ingredient is the symplectic transformations of the quantum canonical coordinates, which are necessary to determine the energy eigenvalues of the quadratic terms. We find the symplectic transformations for the examples with simple classical minima, and further calculate the higher order spectra with standard perturbation methods in quantum mechanics. In Section 4 we systematically compute the periods and topological string free energies for the Calabi-Yau geometries, summarizing the results in previous literature. We then compute the differential operators
which exactly determine quantum corrections to classical periods, generalizing earlier works \[5, 6\] to the situation of higher genus mirror curves. Similarly, the topological free energy in the NS limit is determined by the quantum periods, and we show that this agrees with results from e.g. method of topological vertex. An interesting feature is that the differential operators are the same for differential cycles of the higher genus mirror curves. Following earlier works \[14\], we perform some satisfying tests of our calculations by comparing the quantum spectra from direct perturbation and from Bohr-Sommerfeld quantization using quantum periods as phase volumes. These exercise provide some exact analytic results for the classical and quantum periods of the Calabi-Yau spaces, which are difficult to directly obtain.

2 Dimer models and integrable systems

In \[40\], the authors proposed an infinite class of cluster integrable systems.\(^1\) The most interesting ones among them are the cluster integrable systems for the dimer models on a torus. The dimer model is the study of the set of perfect matching of a graph, where the perfect matching is a subset of edges which covers each vertex exactly once. For a bipartite graph, the vertices are divided into two sets, the black set and the white set. Every edge connects a white vertex to a black vertex. For a more detail introduction to dimer models, see \[45\].

The dimer model can be connected to a toric diagram by Kasteleyn matrix \(K(X, Y)\) \[45\], which is the weighted adjacency matrix of the graph. The determinant of the Kasteleyn matrix, happens to be the mirror curve of the corresponding Toric Calabi-Yau three-fold \[46\][47]. The adjacency matrix can be computed as follows:

- Multiply each edge weight of the graph a sign \(\pm 1\), so that around every face, the product of the edge weights over edges bounding the face is

\[
\text{sgn}(\prod_i e_i) = \begin{cases} +1, & \text{if } (# \text{ edges}) = 2 \mod 4 \\ -1, & \text{if } (# \text{ edges}) = 0 \mod 4 \end{cases}
\]

- Construct two loops \(\gamma_X, \gamma_Y\) along the two cycles of the torus, we draw them as red dash lines in the diagram.

- Fix an orientation, from black to white, as the positive orientation.

- Times each edge with a factor \(X\) or \(Y\), if the loop \(\gamma_X\) or \(\gamma_Y\) get through the edge with positive orientation. Times each edge with a factor \(1/X\) or \(1/Y\), if the loop \(\gamma_X\) or \(\gamma_Y\) get through the edge with positive orientation.

\(^1\)For A type Toda systems, \[43\][44] have an equivalent but different description.
Then the Kasteleyn matrix is a matrix with row labeled by black vertices and column labeled by white vertices, with the entry as the weight between the connected black and white vertices. The entry is 0, if two points are not connected. In this paper, we are interested in $Y^{p,q}$ system, the determinant of the Kasteleyn matrix has the form

$$Y + \frac{X^q}{Y} + X^{p+2} + \cdots + X + 1 = 0 \quad (2.2)$$

Following [40][48], the commutation relations and the Hamiltonians of the cluster integrable systems can be read from the loops of the graph. Let $\omega_i$ be the oriented loops on the graph, the Poisson bracket between cycles are defined as

$$\{\omega_i, \omega_j\} = \epsilon_{\omega_i, \omega_j} \omega_i \omega_j,$$

where

$$\epsilon_{\omega_i, \omega_j} := \sum_v \text{sgn}(v) \delta_v(\omega_i, \omega_j). \quad (2.4)$$

Here sgn$(v) = 1$ for the white vertex $v$, and $-1$ for the black vertex. $\delta_v$ is a skew symmetric bilinear form with $\delta_v(\omega_i, \omega_j) = -\delta_v(\omega_j, \omega_i) = -\delta_v(-\omega_i, \omega_j) \in \frac{1}{2}\mathbb{Z}$, as illustrated in Figure 1. Though more general vertex is possible, for our examples of dimer models we will only encounter cubic vertices.

To construct the basis $t_i$ of all the loops, we can first fix an arbitrary perfect matching as the reference perfect matching, then all the bases can be constructed from the difference between the reference perfect matching and another perfect matching. Then the Hamiltonians $H_n$ are the sum of all possible combinations of $n$ of these cycles $t_i$ with the condition that they do not overlap or touch at any vertex of the tiling.

### 2.1 Examples

In this subsection, we give some examples for the dimer models of $5d, N = 1$ $SU(3)$ gauge theories, with various Chern-Simons level $m = 0, 1, 2, 3$. The graphs of these
theories were appeared during the study of 4d $N = 1$ quiver gauge theories, where
the graphs of the dimer models are brane tiling for the quiver gauge theories. For the $Y^{p,p}$ system, the brane tiling is the well-known Hexagon tiling [48]. One can merge
the points in the tiling for the $Y^{p,p}$ system to get the tiling for a $Y^{p,q}, q < p$ system [49]. For example, the tiling for $Y^{3,3}$ system is depicted in Figure 2a.

$Y^{3,3}$ model

We choose the loops to be\(^2\)

\[
t_1 = 7 \to 1 \to 10 \to 4 \to 7, \quad t_2 = 7 \to 2 \to 11 \to 4 \to 7, \\
t_3 = 8 \to 2 \to 11 \to 5 \to 8, \quad t_4 = 8 \to 3 \to 12 \to 5 \to 8, \\
t_5 = 9 \to 3 \to 12 \to 6 \to 9, \quad t_6 = 9 \to 1 \to 10 \to 6 \to 9.
\]

(2.5)

Only loops that are overlapped have non-vanishing Poisson brackets, they are

\[
\{t_1, t_6\} = -t_1 t_6, \quad \{t_1, t_2\} = t_1 t_2, \quad \{t_2, t_3\} = t_2 t_3, \\
\{t_3, t_4\} = t_3 t_4, \quad \{t_4, t_5\} = t_4 t_5, \quad \{t_5, t_6\} = t_5 t_6.
\]

(2.6)

The Hamiltonians can be read from the graph directly from the rules in precious section:

\[
H_1 = t_1 + t_2 + t_3 + t_4 + t_5 + t_6, \\
H_2 = t_1 t_3 + t_1 t_4 + t_1 t_5 + t_2 t_4 + t_2 t_5 + t_2 t_6 + t_3 t_5 + t_3 t_6 + t_4 t_6, \\
H_3 = t_1 t_3 t_5 + t_2 t_4 t_6.
\]

(2.7)

The Poisson brackets (2.6) can be enhanced to the quantum level as the commutation relations, in terms of canonical variables $\{q_i, p_i = -i\hbar \frac{\partial}{\partial q_i}\}$, we find a possible coordinates relation

\[
t_1 = R^2 e^{q_1}, \quad t_2 = e^{p_1 + q_1}, \quad t_3 = R^2 e^{q_2 - q_1}, \\
t_4 = e^{p_2 + q_2}, \quad t_5 = R^2 e^{-q_2}, \quad t_6 = e^{-p_1 - p_2 - q_1 - q_2}.
\]

(2.8)

The $R$ is the radius of the compactification circle from 5d to 4d, which gives a non-trivial deformation to the integrable systems. It is related to the instanton counting parameter or mass parameters in the 5d gauge theory point of view. One can get the brane tiling of $Y^{3,2}$ systems 2b (b) by merging the point 8, 11 and 2, 5 in Figure 2a. By further merging 9, 12 and 3, 6, we get $Y^{3,1}$ 2c. By doing this further, we get $Y^{3,0}$ 2d. In the following, we list their Poisson brackets and Hamiltonians for these models. The bases $t_i$ are the loops inherited from $t_i$ in $Y^{3,3}$ after merging the points.

\(^2\)There is an independent but irrelevant zig-zag path 1 $\to$ 7 $\to$ 2 $\to$ 8 $\to$ 3 $\to$ 9 $\to$ 1 which commute with other loops we choose. Since it is irrelevant for dynamical Hamiltonians, we don’t mention it in other examples.
Figure 2: Brane tiling for $Y^{3,q}$, $q = 3, 2, 1, 0$, the unit cells are divided by the red dashed lines, which are the loops $\gamma_{X,Y}$ on the torus.
$Y^{3,2}$ model

We choose the loops in Figure 2b

\begin{align}
t_1 &= 6 \rightarrow 1 \rightarrow 9 \rightarrow 4 \rightarrow 6, \quad t_2 = 6 \rightarrow 2 \rightarrow 7 \rightarrow 4 \rightarrow 6, \\
t_3 &= 7 \rightarrow 2 \rightarrow 7, \quad t_4 = 7 \rightarrow 3 \rightarrow 10 \rightarrow 2 \rightarrow 7, \\
t_5 &= 8 \rightarrow 3 \rightarrow 10 \rightarrow 5 \rightarrow 8, \quad t_6 = 8 \rightarrow 1 \rightarrow 9 \rightarrow 5 \rightarrow 8. \\
\end{align}

The non-vanishing Poisson brackets are

\begin{align}
\{t_1, t_6\} &= -t_1 t_6, \quad \{t_1, t_2\} = t_1 t_2, \quad \{t_2, t_3\} = t_2 t_3, \quad \{t_2, t_4\} = t_2 t_4, \\
\{t_3, t_4\} &= t_3 t_4, \quad \{t_4, t_5\} = t_4 t_5, \quad \{t_5, t_6\} = t_5 t_6. \\
\end{align}

In terms of canonical variables,

\begin{align}
t_1 &= R^2 e^{q_1}, \quad t_2 = e^{p_1 + q_1}, \quad t_3 = R^2 e^{q_2 - q_1}, \\
t_4 &= e^{p_2 + q_2 - q_1}, \quad t_5 = R^2 e^{-q_2}, \quad t_6 = e^{-p_1 - p_2 - q_1}. \\
\end{align}

With the Hamiltonians

\begin{align}
H_1 &= t_1 + t_2 + t_3 + t_4 + t_5 + t_6, \\
H_2 &= t_1 t_3 + t_1 t_4 + t_1 t_5 + t_2 t_5 + t_2 t_6 + t_3 t_5 + t_3 t_6 + t_4 t_6, \\
H_3 &= t_1 t_3 t_5. \\
\end{align}

$Y^{3,1}$ model

We choose the loops in Figure 2c

\begin{align}
t_1 &= 5 \rightarrow 1 \rightarrow 8 \rightarrow 4 \rightarrow 5, \quad t_2 = 5 \rightarrow 2 \rightarrow 6 \rightarrow 4 \rightarrow 5, \\
t_3 &= 6 \rightarrow 2 \rightarrow 6, \quad t_4 = 6 \rightarrow 3 \rightarrow 7 \rightarrow 2 \rightarrow 6, \\
t_5 &= 7 \rightarrow 3 \rightarrow 7, \quad t_6 = 7 \rightarrow 1 \rightarrow 8 \rightarrow 3 \rightarrow 7. \\
\end{align}

The non-vanishing Poisson brackets are

\begin{align}
\{t_1, t_6\} &= -t_1 t_6, \quad \{t_1, t_2\} = t_1 t_2, \quad \{t_2, t_3\} = t_2 t_3, \quad \{t_2, t_4\} = t_2 t_4, \\
\{t_3, t_4\} &= t_3 t_4, \quad \{t_4, t_5\} = t_4 t_5, \quad \{t_4, t_6\} = t_4 t_6, \quad \{t_5, t_6\} = t_5 t_6. \\
\end{align}

In terms of canonical variables,

\begin{align}
t_1 &= R^2 e^{q_1}, \quad t_2 = e^{p_1 + q_1}, \quad t_3 = R^2 e^{q_2 - q_1}, \\
t_4 &= e^{p_2 + q_2 - q_1}, \quad t_5 = R^2 e^{-q_2}, \quad t_6 = e^{-p_1 - p_2 - q_1 - q_2}. \\
\end{align}

With the Hamiltonians

\begin{align}
H_1 &= t_1 + t_2 + t_3 + t_4 + t_5 + t_6, \\
H_2 &= t_1 t_3 + t_1 t_4 + t_1 t_5 + t_2 t_5 + t_2 t_6 + t_3 t_5 + t_3 t_6, \\
H_3 &= t_1 t_3 t_5. \\
\end{align}
In terms of canonical variables, Hamiltonians of some orbifold models including $C$ note that the $Y$ for the $Y$ derived from dimer models. In the previous Section 2, we derived the Hamiltonians 0

We choose the loops in Figure 2d

$$t_1 = 4 \rightarrow 1 \rightarrow 4, \quad t_2 = 4 \rightarrow 2 \rightarrow 5 \rightarrow 1 \rightarrow 4,$$

$$t_3 = 5 \rightarrow 2 \rightarrow 5, \quad t_4 = 5 \rightarrow 3 \rightarrow 6 \rightarrow 2 \rightarrow 5,$$

$$t_5 = 6 \rightarrow 3 \rightarrow 6, \quad t_6 = 6 \rightarrow 1 \rightarrow 4 \rightarrow 3 \rightarrow 6.$$  

(2.17)

The non-vanishing Poisson brackets are

$$\{t_1, t_6\} = -t_1 t_6, \quad \{t_1, t_2\} = t_1 t_2, \quad \{t_2, t_3\} = t_2 t_3, \quad \{t_3, t_4\} = t_3 t_4, \quad \{t_4, t_5\} = t_4 t_5,$$

$$\{t_5, t_6\} = t_5 t_6, \quad \{t_2, t_4\} = t_2 t_4, \quad \{t_4, t_6\} = t_4 t_6, \quad \{t_2, t_6\} = -t_2 t_6.$$  

(2.18)

In terms of canonical variables,

$$t_1 = R^2 e^{q_1}, \quad t_2 = e^{p_1 + q_1}, \quad t_3 = R^2 e^{q_2 - q_1},$$

$$t_4 = e^{p_2 - q_1}, \quad t_5 = R^2 e^{-q_2}, \quad t_6 = e^{-p_1 - p_2}.$$  

(2.19)

With the Hamiltonians

$$H_1 = t_1 + t_2 + t_3 + t_4 + t_5 + t_6,$$

$$H_2 = t_1 t_3 + t_1 t_4 + t_1 t_5 + t_2 t_5 + t_3 t_5 + t_3 t_6,$$

$$H_3 = t_1 t_3 t_5.$$  

(2.20)

### 3 Perturbative computations of quantum spectra

In this section, we consider the perturbative energy spectra of the quantum integrable systems described by genus two mirror curves, including the $Y^{3,m}$ models with $m = 0, 1, 2, 3$, and $\mathbb{C}^3/\mathbb{Z}_5$ model. Each model have two dynamical Hamiltonians, which are derived from dimer models. In the previous Section 2, we derived the Hamiltonians for the $Y^{3,m}$ models, where the case of $m = 0$ was also considered in [41]. The Hamiltonians of some orbifold models including $\mathbb{C}^3/\mathbb{Z}_5$ are available in [42]. We also note that the $Y^{3,3}$ model is equivalent to the orbifold $\mathbb{C}^3/\mathbb{Z}_6$ model in [42]. We quantize the Hamiltonians by promoting the dynamical variables to operators with canonical commutation relations $[q_i, q_j] = [p_i, p_j] = 0, [q_i, p_j] = i\hbar \delta_{i,j}$ with $i, j = 1, 2$.

The Hamiltonians are bounded below in the phase space $(q_1, p_1, q_2, p_2)$. First we consider the $Y^{3,0}, Y^{3,3}, \mathbb{C}^3/\mathbb{Z}_5$ models, for which the classical minima are simply located at the origin $q_1 = q_2 = p_1 = p_2 = 0$. We expand the Hamiltonians around the minimal point.

First we study in details the $\mathbb{C}^3/\mathbb{Z}_5$ model, whose Hamiltonians are

$$H_1 = e^{q_1} + e^{p_1} + e^{-q_1 + q_2} + e^{p_2} + e^{-q_2 - p_1 - p_2},$$

$$H_2 = e^{q_2} + e^{q_1 + p_2} + e^{p_1 + p_2} + e^{-p_2 - q_2} + e^{-q_1 - p_1 - p_2}.$$  

(3.1)
We expand the Hamiltonians up to quadratic order

\[ H_i = 5 + \frac{1}{2} (q_1, q_2, p_1, p_2) S_i \begin{pmatrix} q_1 \\ q_2 \\ p_1 \\ p_2 \end{pmatrix} + \mathcal{O}(\hbar^3), \quad i = 1, 2, \quad (3.3) \]

where the \( S_1, S_2 \) are real symmetric matrices

\[
S_1 = \begin{pmatrix} 2 & -1 & 0 & 0 \\ -1 & 2 & 1 & 1 \\ 0 & 1 & 2 & 1 \\ 0 & 1 & 1 & 2 \end{pmatrix}, \quad S_2 = \begin{pmatrix} 2 & 0 & 1 & 2 \\ 0 & 2 & 0 & 1 \\ 1 & 0 & 2 & 2 \\ 2 & 1 & 2 & 4 \end{pmatrix}. \quad (3.4)
\]

We would like to write the quadratic Hamiltonians as linear combinations of two harmonic oscillators. We consider a linear transformation

\[
\begin{pmatrix} q_1 \\ q_2 \\ p_1 \\ p_2 \end{pmatrix} = M \begin{pmatrix} x_1 \\ x_2 \\ y_1 \\ y_2 \end{pmatrix}, \quad (3.5)
\]

where \( M \) is a \( 4 \times 4 \) real matrix. To preserve the same canonical commutation relation, the matrix \( M \) must be a symplectic matrix \( M \Sigma M^T = \Sigma \), where \( \Sigma \) is the antisymmetric matrix

\[
\Sigma = \begin{pmatrix} 0 & 0 & 1 & 0 \\ 0 & 0 & 0 & 1 \\ -1 & 0 & 0 & 0 \\ 0 & -1 & 0 & 0 \end{pmatrix}. \quad (3.6)
\]

It turns out due to the special property that the Hamiltonians commute with each other, we can find symplectic transformation \( M \) so that the quadratic terms can be written as linear combinations of the two harmonic oscillators

\[
H_1 = 5 + \frac{1}{2} [c_1(x_1^2 + y_1^2) + c_2(x_2^2 + y_2^2)] + \mathcal{O}(\hbar^{\frac{3}{2}}),
\]

\[
H_2 = 5 + \frac{1}{2} [c_3(x_1^2 + y_1^2) + c_4(x_2^2 + y_2^2)] + \mathcal{O}(\hbar^{\frac{3}{2}}). \quad (3.7)
\]

There is a continuous 2-parameter family of solutions for the matrix \( M \). Without loss of generality, we can use a particular solution

\[
M_{\mathbb{C}^3/\mathbb{Z}_5} = \begin{pmatrix}
-\frac{(5-2\sqrt{5})^{\frac{1}{2}}}{\sqrt{\frac{4}{5}}} & -\frac{(10-2\sqrt{5})^{\frac{1}{2}}}{\sqrt{\frac{3}{5}}} & (\frac{1}{4} + \frac{1}{2\sqrt{5}})^{\frac{1}{4}} & 0 \\
-\frac{\sqrt{2}}{2}(3-2\sqrt{5})^{\frac{1}{2}} & -\frac{6}{2^{\frac{1}{4}}(1 + \frac{1}{\sqrt{5}})} & 0 & -\frac{1}{2^{\frac{1}{4}}(1 - 2\sqrt{5})^{\frac{1}{4}}} \\
-\frac{(10+2\sqrt{5})^{\frac{1}{2}}}{\sqrt{\frac{2}{5}}} & (\frac{1}{2^\frac{1}{4}} + \frac{1}{10\sqrt{5}})^{\frac{1}{4}} & 0 & -\frac{1}{2^{\frac{1}{4}}(1 - 2\sqrt{5})^{\frac{1}{4}}} \\
(\frac{1}{8} + \frac{11}{40\sqrt{5}})^{\frac{1}{4}} & -\frac{(25-11\sqrt{5})^{\frac{1}{2}}}{2^{\frac{1}{4}}\sqrt{\frac{3}{5}}} & -\frac{1}{2^{\frac{1}{4}}(1 - \sqrt{5})^{\frac{1}{4}}} & -\frac{1}{2^{\frac{1}{4}}(1 + \sqrt{5})^{\frac{1}{4}}} \\
\end{pmatrix}, \quad (3.8)
\]
with the linear coefficients
\[ c_1 = \left(\frac{5 + \sqrt{5}}{2}\right)^\frac{1}{2}, \quad c_2 = \left(\frac{5 - \sqrt{5}}{2}\right)^\frac{1}{2}, \]
\[ c_3 = (5 - 2\sqrt{5})^\frac{1}{2}, \quad c_4 = (5 + 2\sqrt{5})^\frac{1}{2}. \]  
(3.9)

Denoting the quantum levels of the harmonic oscillators \((x_1, y_1)\) and \((x_2, y_2)\) by two non-negative integers \(n_1, n_2\), the quantum spectrum up to order \(\hbar\) is
\[ \left(\frac{E_1}{E_2}\right) = 5 \left(\frac{1}{1}\right) + \left(\frac{c_1}{c_3} \frac{c_2}{c_4}\right) \left(\frac{n_1 + \frac{1}{2}}{n_2 + \frac{1}{2}}\right) \hbar + \mathcal{O}(\hbar^2) \]  
(3.10)

We can further compute the higher order corrections to energy spectra. We use the time-independent perturbation theory well-known in quantum mechanics, which separates a Hamiltonian into a zero order part and a perturbation part
\[ \mathcal{H} = \mathcal{H}_0 + \mathcal{H}', \]  
(3.11)

where the zero order part \(\mathcal{H}_0\) corresponds to the Hamiltonians up to quadratic order in (3.3), while the perturbation part \(\mathcal{H}'\) corresponds to the higher order terms.

We denote the harmonic quantum states of the zero order Hamiltonians as \(|n_1, n_2\rangle\).

Then the first few order corrections to energy spectra are
\[ E_{(n_1, n_2)} = E_{(n_1, n_2)}^{(0)} + \langle n_1, n_2 | \mathcal{H}' | n_1, n_2 \rangle 
+ \sum_{(m_1, m_2) \neq (n_1, n_2)} \frac{\langle m_1, m_2 | \mathcal{H}' | n_1, n_2 \rangle^2}{E_{(n_1, n_2)}^{(0)} - E_{(m_1, m_2)}^{(0)}} + \cdots. \]  
(3.12)

To compute the next \(\hbar^2\) order corrections, we need to expand the exponentials in the Hamiltonians (3.1) to cubic and quartic orders, and rewrite the canonical coordinates in terms of the standard creation and annihilation operators. For the first correction \(\langle n_1, n_2 | \mathcal{H}' | n_1, n_2 \rangle\), the cubic terms have no contribution since there is an odd number of creation and annihilation operators, while the quartic terms make an order \(\hbar^2\) contribution. The cubic terms have a \(\hbar^2\) order contribution in the more complicated second correction term in the above equation (3.12). After some complicated calculations, we find the \(\hbar^2\) order contributions to the quantum spectra. For the \(\mathbb{C}^3/\mathbb{Z}_5\) model, the results are
\[ E_1 = 5 + \left[\left(\frac{5 + \sqrt{5}}{2}\right)^\frac{1}{2} n_1 + \left(\frac{5 - \sqrt{5}}{2}\right)^\frac{1}{2} n_2 + \frac{1}{2}(5 + 2\sqrt{5})^\frac{1}{2}\right] \hbar 
+ [7 + 2(3 + \sqrt{5})n_1(1 + n_1) + 2(3 - \sqrt{5})n_2(1 + n_2)] \frac{\hbar^2}{40} + \mathcal{O}(\hbar^3), \]  
(3.13)

\[ E_2 = 5 + \left[\left(5 - 2\sqrt{5}\right)^\frac{1}{2} n_1 + \left(5 + 2\sqrt{5}\right)^\frac{1}{2} n_2 + \left(\frac{5 + \sqrt{5}}{2}\right)^\frac{1}{2}\right] \hbar 
+ \left[3 + \sqrt{5} + 4n_1 
+ 2(2 - \sqrt{5})n_1^2 + 4(1 + \sqrt{5})n_2 + 2(2 + \sqrt{5})n_2^2 + 4\sqrt{5}n_1n_2\right] \frac{\hbar^2}{20} + \mathcal{O}(\hbar^3). \]
It is well known that the eigenvalues of a matrix do not change under a similarity transformation of the matrix. Here analogously we find that the spectra in (3.13) are independent of the choice of symplectic transformation, up to the trivial freedom of exchanging the two quantum numbers $n_1 \leftrightarrow n_2$. This is easy to understand from a physics point of view since the Hamiltonians are the same regardless of the choices of the canonical coordinates. Furthermore, the linear coefficients (3.9) are indeed related to the eigenvalues of certain matrices. We note that for a general even-dimensional real symmetric matrix $S$, since $\det(S\Sigma - \lambda I) = \det(S\Sigma + \lambda I) = \det(S\Sigma + \lambda I)$, the eigenvalues of $S\Sigma$ and $\Sigma$ are the same and always come in pairs with opposite signs. In our context, we find that for the matrices (3.4) in the quadratic Hamiltonians, the eigenvalues of $S\Sigma$ and $\Sigma$ are always purely imaginary and the positive imaginary parts are exactly the linear coefficients (3.9). Namely, the eigenvalues of $S\Sigma$ are $\pm i c_1, \pm i c_2$ and the eigenvalues of $S\Sigma$ are $\pm i c_3, \pm i c_4$. This is also true for the $Y^{3,0}$ and $Y^{3,3}$ models discussed below. In the Appendix A we give a simple general mathematical proof of this property.

Similarly we find the symplectic transformations and the perturbative energy spectra for the $Y^{3,0}$ and $Y^{3,3}$ models in (2.20, 2.7). Again there is a continuous 2-parameter family of solutions for symplectic transformations. For the $Y^{3,0}$ model, we can use for example a solution

$$M_{Y^{3,0}} = \frac{1}{\sqrt{2R^3}} \left( \begin{array}{cccc} -R & -2R & -\sqrt{3}R & 0 \\ -2R & -R & 0 & -\sqrt{3}R \\ \sqrt{R^2 + 4} - R & 0 & -\sqrt{3}R & \sqrt{3}R \\ \frac{\sqrt{R^2 + 4} + R}{3R - \sqrt{R^2 + 4}} & 0 & 0 & 0 \end{array} \right),$$

and the perturbative energy spectrum is

$$\begin{pmatrix} E_1 \\ E_2 \end{pmatrix} = 3(1 + R^2) \left( \begin{array}{c} 1 \\ 1 \end{array} \right) + \frac{\sqrt{3}R}{2} \left[ \sqrt{4 + R^2}(n_1 + n_2 + 1) \left( \begin{array}{c} 1 \\ 1 \end{array} \right) + R(n_1 - n_2) \left( \begin{array}{c} 1 \\ -1 \end{array} \right) \right] \hbar$$

$$\left\{ \left[ 4(11 + 15n_1 + 6n_1^2 + 15n_2 + 6n_2^2 + 18n_1n_2) + 5 + 6(n_1 + n_1^2 + n_2 + n_2^2) \right] R^2$$

$$- \frac{72}{4 + R^2}(2 + 3n_1 + n_1^2 + n_2 + n_2^2 + 4n_1n_2) \left( \begin{array}{c} 1 \\ 1 \end{array} \right) \right\} \frac{\hbar^2}{72} + O(\hbar^3). \quad (3.14)$$

We see there is an apparent symmetry of the spectra. The spectra of the two Hamiltonians $E_1 \leftrightarrow E_2$ are exchanged if the quantum levels are exchanged $n_1 \leftrightarrow n_2$.

For the $Y^{3,3}$ model, the results are

$$M_{Y^{3,3}} = 2 \cdot 3^{\frac{3}{2}} \sqrt{R} \left( \begin{array}{cccc} -R & -2R & -\sqrt{3}R & 0 \\ -2R & -R & 0 & -\sqrt{3}R \\ R + 2 & 2R - 1 & \sqrt{3}R & -\sqrt{3} \\ 2R - 1 & R + 2 & -\sqrt{3} & \sqrt{3}R \end{array} \right),$$

11
The first Hamiltonian of the perturbation in the degenerate space actually vanish, i.e.
\[ \langle \, \rangle = 0 \] for quantum states with the same classical minima by solving for the critical points of Hamiltonians
\[ \partial_i H = 0 \] for \( i = 1, 2 \). We find that the minima are located at the same points for the two Hamiltonians of the quantum system due to the special property that the Hamiltonians commute with each other. In these models it is much more complicated to find the symplectic transformations that diagonalize the quadratic terms of the Hamiltonians of the quantum system due to the special property that the Hamiltonians commute with each other. In these models it is much more complicated to find the symplectic transformations that diagonalize the quadratic terms of the Hamiltonians expanded around the minima. However, we can still use the formula in Appendix A to compute the \( \hbar \) order contributions to quantum spectra in terms of the eigenvalues of certain matrices from the quadratic terms.

For the remaining \( Y^{3,1} \) and \( Y^{3,2} \) models (2.16, 2.12), we need to determine the classical minima by solving for the critical points of Hamiltonians \( \partial_q H = \partial_p H = 0 \) for \( i = 1, 2 \). We find that the minima are located at the same points for the two Hamiltonians of the quantum system due to the special property that the Hamiltonians commute with each other. In these models it is much more complicated to find the symplectic transformations that diagonalize the quadratic terms of the Hamiltonians expanded around the minima. However, we can still use the formula in Appendix A to compute the \( \hbar \) order contributions to quantum spectra in terms of the eigenvalues of certain matrices from the quadratic terms.

For the \( Y^{3,1} \) model, the minima are at
\[ q_1 = -3 \log(r), \quad q_2 = -6 \log(r), \quad p_1 = - \log(r), \quad p_2 = - \log(r + R^2), \] (3.16)
where \( r \) is the only positive root of the polynomial equation,
\[ r^9 + R^2 r^8 = 1, \] (3.17)
with numerical value e.g. \( r = 0.921599 \) for \( R = 1 \). The quantum spectra are
\[ E_1 = \frac{3r + 4R^2}{r^6(r + R^2)} + \frac{r^2 R[(3r + 2R^2 + 2R\sqrt{r + R^2})^\frac{1}{2}]}{r^6(r + R^2)}(n_1 + \frac{1}{2}) \]
\[ + (3r + 2R^2 - 2R\sqrt{r + R^2})^\frac{1}{2}(n_2 + \frac{1}{2})] \hbar + \mathcal{O}(\hbar^2), \] (3.18)
\[ E_2 = \frac{3r + 2R^2}{r^4} + \frac{r^2 R(r + R^2)^\frac{1}{2}}{r^4}[(3r + 4R^2 - 4R\sqrt{r + R^2})^\frac{1}{2}](n_1 + \frac{1}{2}) \]
\[ + (3r + 4R^2 + 4R\sqrt{r + R^2})^\frac{1}{2}(n_2 + \frac{1}{2})] \hbar + \mathcal{O}(\hbar^2). \]
For the $Y^{3.2}$ model, the minima are at
\[ q_1 = -3 \log[r(r + R^2)], \quad q_2 = -\log[r^2(r + R^2)], \]
\[ p_1 = -\log(r), \quad p_2 = -\log(r^2), \] (3.19)
where $r$ is now the only positive root of the equation
\[ r^9 + 4R^2r^8 + 6R^4r^7 + 4R^6r^6 + R^8r^5 = 1, \] (3.20)
with numerical value e.g. $r = 0.665055$ for $R = 1$. The quantum spectra are
\[ E_1 = \frac{3r + 5R^2}{r^7(r + R^2)^6} + \frac{r^3R(r + R^2)^2}{\sqrt{2}} [(6r + 5R^2 + R\sqrt{4r + 5R^2})^\frac{1}{2}(n_1 + \frac{1}{2}) \]
\[ + (6r + 5R^2 - R\sqrt{4r + 5R^2})^\frac{1}{2}(n_2 + \frac{1}{2})]h + O(h^2), \] (3.21)
\[ E_2 = \frac{3r^2 + 7rR^2 + 5R^4}{r^4(r + R^2)^4} + \frac{r^2R(r + R^2)^2}{\sqrt{2}} \{[r^3(r + R^2)^2(6r^3 + 25r^2R^2 + 30rR^4 + 10R^6) \]
\[ - R(5r^2 + 10rR^2 + 4R^4)\sqrt{r(4r + 5R^2)}]^\frac{1}{2}(n_1 + \frac{1}{2}) + [r^3(r + R^2)^2(6r^3 + 25r^2R^2) \]
\[ + 30rR^4 + 10R^6) + R(5r^2 + 10rR^2 + 4R^4)\sqrt{r(4r + 5R^2)}]^\frac{1}{2}(n_2 + \frac{1}{2})\}h + O(h^2). \]
Without solving the symplectic transformations for these two $Y^{3.1}$ and $Y^{3.2}$ models, there is an ambiguity of exchanging the quantum numbers $n_1 \leftrightarrow n_2$ in the spectra. This can be fixed by comparing with the derivatives of periods of the corresponding Calabi-Yau geometries.

4 From topological strings to energy spectra

In this section we will show that the spectrum problem can be solved by utilizing the well-known method in topological string theory. More precisely, we use the method developed in [4], and calculate the energy spectra by imposing the Bohr–Sommerfeld quantization condition on the quantum B-periods of the mirror curves. First we summarize some basic facts about the classical/quantum mirror curves, and the general relations between topological strings and the energy spectra. After that, we will demonstrate how we calculate the energy spectra from the quantum periods in some concrete models.

4.1 General aspects of classical/quantum curves

We consider topological string theory on the toric Calabi-Yau three-fold, where the topological information in the B model are captured by a mirror curve. A genus-$g$ mirror curve is defined by the algebraic equation for $x, y \in \mathbb{C},$
\[ W(e^x, e^y; z) = 0, \] (4.1)
where $z = (z_1, z_2, ..., z_s)$ are the complex structure moduli parameters with $s := b^{1,1} \geq g$. Generally, there are $g$ dynamical moduli corresponding to $g$ compact A- and B-cycles of the Riemann surface, and the $s - g$ remaining ones are known as non-dynamical mass parameters. We can define two kinds of classical periods called as A- and B-periods by integrating $y = y(x; z)$ around compact A-cycles and their dual B-cycles,

$$
\Pi_i(z) = \oint_{A_i} y(x; z) dx, \quad \Pi_{i,d}(z) = \oint_{B_i} y(x; z) dx, \quad (i = 1, ..., g)
$$

(4.2)

where $y(x; z)$ is the solution of (4.1).

The mirror maps connecting the Kähler moduli with the complex moduli can be written as linear combinations of A-periods and mass parameters

$$
t_j(z) = \sum_{i=1}^{g} C_{ij} \Pi_i(z) + \text{mass terms}, \quad j = 1, 2, \cdots, s,
$$

(4.3)

where the mass terms depend only on logarithm of mass parameters and will not appear in quantum corrections. Here $C_{ij}$ is the intersection matrix of compact divisors and the base curves we have chosen. With a suitable choice of base curves, parts of the $g \times s$ matrix $C_{ij}$ happens to be the Cartan matrix of the gauge group in the context of geometric realizations of gauge theories.

In the similar way, the dual B-periods give the derivatives of the genus-zero topological string amplitude, so-called prepotential $F_0(t)$,

$$
\Pi_{d,i}(z) = \frac{\partial F_0(t(z))}{\partial \Pi_i(z)} = \sum_{j=1}^{s} C_{ij} \frac{\partial F_0(t(z))}{\partial t_j(z)}, \quad i = 1, 2, \cdots, g.
$$

(4.4)

From the prepotential, we define the Bohr-Sommerfeld volumes as the derivatives of prepotential with appropriate shift $4\pi^2 b^{NS}_i$

$$
\text{vol}_i(z) = \sum_{j=1}^{s} C_{ij} \left( \frac{\partial F_0(t)}{\partial t_j} + 4\pi^2 b^{NS}_j \right).
$$

(4.5)

This shift can be derived from the S-dual like invariance of the classical volumes [50]. It can be absorbed into the genus zero free energy by adding a $t_i$ linear term [22]. In gauge theory point of view, $b^{NS}_i$ comes from the one loop contribution. For 5d $\mathcal{N} = 1$ pure $SU(N)$ gauge theories with Chern-Simons level, denoting the $t_i$, $i \leq N - 1$ the node of $A_N$ group, and $t_N$ the instanton counting parameter. By setting $b^{NS}_N = 0$, we have $b^{NS}_i = b^{NS}_{N-i} = -\frac{(N-i)}{12}$ for $i = 1, 2, \cdots, \frac{N}{2}$. For the $SU(3)$ models we considered, we always have $b^{NS}_1 = b^{NS}_2 = -\frac{1}{6}$, $b^{NS}_3 = 0.$
For a toric Calabi-Yau three-fold, an efficient way to calculate A- and B-period is to solve the Picard–Fuchs equations defined by

\[ \mathcal{L}_\alpha \Pi_i = 0, \quad \mathcal{L}_\alpha \Pi_{d,i} = 0, \]

\[ \mathcal{L}_\alpha = \left[ \prod_{Q_i > 0} \left( \frac{\partial}{\partial x_i} \right)^{Q_i^\alpha} - \prod_{Q_i > 0} \left( \frac{\partial}{\partial x_i} \right)^{Q_i^\alpha} \right], \quad (4.6) \]

where \( Q_i^\alpha \) is the charge vector and \( x_i \) is the homogeneous coordinate of the toric variety. The differential operator \( \mathcal{L}_\alpha \)'s are known as the Picard–Fuchs operators. The variables \( x_i \) relate to \( z \) through the Batyrev coordinates

\[ z_\alpha = \prod_{i=1}^{k+3} x_i^{Q_i^\alpha}. \quad (4.7) \]

The A- and B-periods correspond to logarithmic and double-logarithmic solutions.

Now we promote the classical variables \( x, y \) to the quantum operators \( x, y \) with the canonical commutation relation,

\[ [x, y] = i\hbar. \quad (4.8) \]

Accordingly, the mirror curve is replaced by the difference equation,

\[ W(e^x, e^y)\Psi(x) = 0, \quad (4.9) \]

where \( \Psi(x) \) is the wave function. We can solve the difference equation by utilizing the WKB analysis,

\[ \Psi(x) = \exp \left( \frac{i}{\hbar} \int x w(x'; \hbar)dx' \right). \quad (4.10) \]

Then, we can define quantum version of two periods, called as quantum A- and B-period,

\[ \Pi_i(z) \rightarrow \Pi_i(z; \hbar) = \sum_{n=0}^{\infty} \Pi_i^{(n)} \hbar^n, \quad \Pi_i^{(n)} = \oint_A w^{(n)}(x)dx, \quad (4.11a) \]

\[ \Pi_{d,i}(z) \rightarrow \Pi_{d,i}(z; \hbar) = \sum_{n=0}^{\infty} \Pi_{i,d}^{(n)} \hbar^n, \quad \Pi_{i,d}^{(n)} = \oint_B w^{(n)}(x)dx, \quad (4.11b) \]

where we expand \( w(x; \hbar) \) in \( \hbar \),

\[ w(x; \hbar) = \sum_{n=0}^{\infty} w^{(n)} \hbar^n. \quad (4.12) \]

In our example, \( w^{(2n-1)} \), \( n \in \mathbb{Z}_{>0} \) can be expressed as the total derivative of simple functions with no monodromy. Thus, its contour integral vanishes, and only \( \hbar^{2n} \)-corrections survive.
The quantum corrected prepotential $F(t; \hbar)$, so-called NS free energy, is defined by refined topological string free energy in the NS limit

$$F(t, \hbar) = \sum_{n=0}^{\infty} F_n(t)\hbar^{2n}. \quad (4.13)$$

It satisfies a similar equation as the prepotential

$$\Pi_{d,i}(z; \hbar) = \sum_{j=1}^{s} C_{ij} \frac{\partial F(t(z; \hbar), \hbar)}{\partial t_j(z, \hbar)}, \quad (4.14)$$

where $t_i(z; \hbar)$ are the quantum mirror maps. Comparing both sides of (4.14), we can obtain the recursion relations which enable us to fix $F_i(t)$ completely, up to irrelevant constants and mass parameters.

The Bohr–Sommerfeld volumes (4.5) also have quantum corrections,

$$\text{vol}_i(z) \rightarrow \text{vol}_i(z; \hbar) = \sum_{n \geq 0} \text{vol}_i^{(2n)}(z)\hbar^{2n}. \quad (4.15)$$

In quantum mechanics, the phase volume should be quantized. In our case, the B-periods are quantized,

$$\oint_{B_i} w(x, \hbar)dx = 2\pi\hbar \left( n_i + \frac{1}{2} \right). \quad n_i \in \mathbb{Z}_{\geq 0}. \quad (4.16)$$

By using (4.5), we can rewrite the quantization condition as follows,

$$\text{vol}_i(z; \hbar) = 2\pi\hbar \left( n_i + \frac{1}{2} \right), \quad i = 1, 2, \ldots, g. \quad (4.17)$$

The dynamical complex structure moduli will correspond to Hamiltonians of the quantum systems as we will see in concrete examples. As in the case of NS free energy, by expanding the quantum B-periods in $\hbar$, we can determine the quantum corrections to the energy eigenvalues recursively. The B-periods have to vanish in the classical limit at the minimal energy points, which correspond to the conifold points in the topological string moduli space. Thus, to solve the spectral problem from the topological strings, we have to calculate the phase volumes at the conifold point. It turns out that there is no logarithmic cut for the classical volumes (B-periods) at the conifold points, so they are the same as the mirror maps up to numerical factors,

$$\text{vol}_i(\text{Coni}; \hbar) \sim t_{i,c}(\text{Coni}; \hbar), \quad (4.18)$$

where Coni denotes the conifold point, and $t_{i,c}(z_c; \hbar)$ is quantum mirror map expanded around the conifold point. The numerical factor in the coefficients of $t_{i,c}(z_c; \hbar)$ can be determined by comparing with the derivatives of the classical volumes at conifold point or the perturbative computation as we have done in the previous section.
Therefore, we can calculate the eigenvalues only by using the quantum mirror maps near the conifold points.

Now we move to the computation of the quantum periods. It is straightforward to calculate the quantum A-periods from the definition by taking residues, whereas the direct computations of B-periods are usually not so easy. Here we utilize the differential operator method proposed in [3], and developed in [5].

The important fact is that the quantum A-periods can be given by the classical periods with differential operators as follows,

\[ \Pi_k(z; \hbar) = \left( \sum_{n=0}^{\infty} \hbar^{2n} D_{2n} \right) \Pi_k(z), \quad k = 1, 2, \ldots, g, \tag{4.19} \]

where

\[ D_{2n} = D_{2n}(\theta_{z_1}, \theta_{z_2}, \ldots \theta_{z_s}), \quad \theta_{z_i} = z_i \frac{\partial}{\partial z_i}. \tag{4.20} \]

and coefficients of \( \theta_{z_i} \) are given by rational functions of \( z_i \). This means that we can obtain the differential operators in the conifold frame by transforming from large radius frame to the conifold frame \( z_i \rightarrow z_{c,i} \). Then, by acting the operators on the classical A-periods expanded near the conifold point, we can obtain the quantum corrections in the conifold frame. Since the mass parameters are annihilated by the differential operators, they do not receive quantum corrections.

According to (4.3), the quantum mirror maps are determined by the same differential operators as

\[ t_i(z; \hbar) = \left( \sum_{n=0}^{\infty} \hbar^{2n} D_{2n} \right) t_i(z), \quad i = 1, 2, \ldots, s. \tag{4.21} \]

Interestingly, the differential operators that we will treat in our study do not depend on the choice of the cycles\(^3\). Also, the classical mirror maps can be calculated from the Picard–Fuchs operators. Therefore, it is enough to calculate one of the quantum A-periods to derive the differential operators and determine the quantum mirror maps.

By combining (4.18) with (4.21), the quantum corrections to the volumes and their derivatives with respect to the eigenvalues are given by

\[ \partial^p_{E_1} \partial^p_{E_2} \cdots \partial^p_{E_s} \text{vol}^{(2n)}_j \sim \partial^p_{E_1} \partial^p_{E_2} \cdots \partial^p_{E_s} (D_{2n} t_{c,j}), \tag{4.22} \]

where \( p_i \in \mathbb{Z}_{\geq 0}, n \in \mathbb{Z}_{>0} \), and \( j = 1, 2, \ldots, s \). To calculate the right hand side, we use \( \partial_{E_i} = \sum_{j=1}^{s} (\partial_{E_i z_{c,j}}) \partial_{z_{c,j}} \).

Remarkably, this structure is the same as the quantum B-period; the quantum corrections can be calculated by acting above operators on the classical B-periods,

\[ \Pi_{d,i}(z; \hbar) = \left( \sum_{n=0}^{\infty} \hbar^{2n} D_{2n} \right) \Pi_{d,i}(z). \tag{4.23} \]

\(^3\)It would be interesting to confirm this property in general setup.
This means that once we derive the differential operators $\mathcal{D}_{2n}$ from the quantum A-period that we know how to calculate systematically, we can obtain the quantum B-period which is difficult to obtain by the direct computation of the cycle integral.

Similar to previous paper [5], we can derive recursion relations for the NS free energy by expanding the equations (4.21, 4.14, 4.13). We can explicitly do this for the first and second correction terms $F_1, F_2(t)$, which are determined by the differential operators $\mathcal{D}_2, \mathcal{D}_4$. In our examples, the differential operators will be a linear combinations of first and second derivatives of the complex structure moduli. Suppose

$$\mathcal{D}_2 = \sum_i s_i^{(2)} \theta_i + \sum_{i,j} s_{i,j}^{(2)} \theta_i \theta_j, \quad (4.24)$$

where the coefficients $s_i, s_{i,j}$ are rational functions of complex structure moduli $z_i$’s. Denote the classical mirror maps as $t_i$, then it is straightforward to compute

$$\theta_k(\partial_{t_i} F_0) = \sum_j \theta_k(t_j)(\partial_{t_i} \partial_{t_j} F_0), \quad (4.25)$$

$$\theta_k \theta_l(\partial_{t_i} F_0) = \sum_j \theta_k(t_j)(\partial_{t_i} \partial_{t_j} F_0) + \sum_{j,m} \theta_k(t_j) \theta_l(t_m)(\partial_{t_i} \partial_{t_j} \partial_{t_m} F_0).$$

So we have

$$\mathcal{D}_2(\partial_{t_i} F_0) = \sum_j \mathcal{D}_2(t_j)(\partial_{t_i} \partial_{t_j} F_0) + \sum_{j,k,l,m} s_{l,m}^{(2)} \theta_l(t_j) \theta_m(t_k)(\partial_{t_i} \partial_{t_j} \partial_{t_k} F_0). \quad (4.26)$$

Combining the $\hbar^2$ equations of (4.13, 4.14, 4.21), we find the linear coefficients $s_i^{(2)}$ cancel out. The equation for first order NS free energy is then

$$\sum_{i=1}^s C_{n,i}[\partial_{t_i} F_1 - \sum_{j,k,l,m} s_{l,m}^{(2)} \theta_l(t_j) \theta_m(t_k)(\partial_{t_i} \partial_{t_j} \partial_{t_k} F_0)] = 0, \quad n = 1, 2, \cdots, g. \quad (4.27)$$

If $s = g$ and the matrix $C_{i,j}$ is invertible, it cancels out in the above equation. Otherwise, in general we need to solve the equations including the $C$ matrix. Similarly, repeating the same computation to the next order, we have

$$\sum_{i=1}^s C_{n,i}[\partial_{t_i} F_2 - \sum_{j,k,l,m} s_{l,m}^{(4)} \theta_l(t_j) \theta_m(t_k)(\partial_{t_i} \partial_{t_j} \partial_{t_k} F_0) - \sum_j \mathcal{D}_2(t_j)(\partial_{t_i} \partial_{t_j} F_1)$$

$$- \frac{1}{2} \sum_{j,k} \mathcal{D}_2(t_j) \mathcal{D}_2(t_k)(\partial_{t_i} \partial_{t_j} \partial_{t_k} F_0)] = 0, \quad n = 1, 2, \cdots, g. \quad (4.28)$$

Again, the linear coefficients $s_i^{(4)}$ cancel out. By using (4.27), if the matrix $C$ is invertible, we can eliminate $F_1$, and obtain the relation between $F_2$ and $F_0$. 

18
4.2 Examples

We shall demonstrate the previous computations in some concrete models. In our examples, we focus on the genus-2 mirror curves: $\mathbb{C}^3/\mathbb{Z}_5$ and $Y^{3,m}$ with $m = 0, 1, 2, 3$. Most of classical computations have already done in e.g. [36, 51, 41, 42, 22], and we gather the results to make the paper self-contained. In the following computations, we may omit some arguments in the functions for short notation.

4.2.1 $\mathbb{C}^3/\mathbb{Z}_5$ case

The mirror curve of $\mathbb{C}^3/\mathbb{Z}_5$ is defined as
\[
e^x + e^{-x-p} + e^{-p} + z_1^{1/3} z_2 e^{2x} + z_1^{-1/3} = 0. \tag{4.29}
\]
The Picard–Fuchs operators are
\[
\mathcal{L}_1 = -2\theta_1^2 \theta_2 + \theta_1^3 + z_1(-2\theta_2 + 3\theta_2^2 - \theta_2^3 + 3\theta_2 - 3\theta_1 \theta_2 + 9\theta_1 \theta_2^2 + 27\theta_2^3 - 27\theta_1^2 \theta_2 + 27\theta_1^3),
\]
\[
\mathcal{L}_2 = \theta_2^2 - 3\theta_1 \theta_2 + z_2(-2\theta_2 - 4\theta_2^2 + \theta_1 + 4\theta_1 \theta_2 - \theta_2^3),
\]
\[
\mathcal{L}_3 = \theta_1^2 \theta_2 + z_1 z_2(-2\theta_2^2 + 2\theta_2^3 + 7\theta_1 \theta_2 - 13\theta_1 \theta_2^2 - 3\theta_2^3 + 24\theta_1^2 \theta_2 - 9\theta_1^3). \tag{4.30}
\]

To provide the solutions of the Picard–Fuchs equations, we first define following function,
\[
\omega_0(\rho_i) = \sum_{l,m \geq 0} c(l, m; \rho) z_1^{l+\rho_1} z_2^{m+\rho_2} \tag{4.31}
\]
with
\[
c(l, m; \rho) = \frac{\Gamma(p_1 + 1)^2 \Gamma(p_2 + 1) \Gamma(p_1 - 2p_2 + 1) \Gamma(-3p_1 + p_2 + 1)}{\Gamma(l + p_1 + 1)^2 \Gamma(m + p_2 + 1) \Gamma(l - 2m + p_1 - 2p_2 + 1) \Gamma(-3l + m - 3p_1 + p_2 + 1)}. \tag{4.32}
\]
We further define the derivatives of $\omega_0(\rho_i)$,
\[
\omega_i = \left. \frac{\partial \omega_0}{\partial \rho_i} \right|_{\rho_{1,2}=0}, \quad \omega_{ij} = \left. \frac{\partial^2 \omega_0}{\partial \rho_i \partial \rho_j} \right|_{\rho_{1,2}=0}. \tag{4.33}
\]
Then, the mirror maps are given by,
\[
t_1(z) = \omega_1 = \log z_1 - 6z_1 - z_2 + 45z_1^2 - \frac{3}{2} z_2^2 + \mathcal{O}(z_1^3),
\]
\[
t_2(z) = \omega_2 = \log z_2 + 2z_1 + 2z_2 - 15z_1^2 + 3z_2^2 + \mathcal{O}(z_1^3). \tag{4.34}
\]
The derivatives of the prepotential are
\[
\frac{\partial F_0}{\partial t_1} = 2\omega_{1,1} + 2\omega_{1,2} + 3\omega_{2,2},
\]
\[
\frac{\partial F_0}{\partial t_2} = \omega_{1,1} + 6\omega_{1,2} + 9\omega_{2,2}. \tag{4.35}
\]
The classical B-periods $\Pi_{d,i}$ ($i = 1, 2$) are given by the formula (4.4), where the matrix $C_{ij}$ of this model is,

$$C = \begin{bmatrix} 3 & -1 \\ -1 & 2 \end{bmatrix}. \quad (4.36)$$

From the prepotential, the Bohr-Sommerfeld volumes are

$$\text{vol}^{(0)}_1(z) = 3 \frac{\partial F_0}{\partial t_1} - \frac{\pi^2}{2},$$
$$\text{vol}^{(0)}_2(z) = -\frac{\partial F_0}{\partial t_2} + 2 \frac{\partial F_0}{\partial t_2} - \frac{2\pi^2}{3}. \quad (4.37)$$

where the complex structure moduli $z_1, z_2$ are related to the quantum systems of the dimer model by

$$z_1 = -\frac{E_1}{E_2}, \quad z_2 = \frac{E_2}{E_1}. \quad (4.38)$$

The classical volumes vanish at conifold point, $z_1 = -1/25$, $z_2 = 1/5$, or $E_1 = E_2 = 5$, which is checked numerically.

Now let us consider the quantum periods. Correspondingly, the classical mirror curve is replaced by the difference equation,

$$\Psi(x + i\hbar) + e^{-\frac{i\theta_1}{2}} \Psi(x - i\hbar) + \left( z_1^{\frac{3}{2}} z_2 e^{2x} + e^{x} + z_1^{-\frac{1}{2}} \right) \Psi(x) = 0. \quad (4.39)$$

According to [4], the quantum A-periods are given by taking the residue,

$$\Pi(z; \hbar) = \frac{1}{5} \log(z_1^2 z_2) + \int_{x=-\infty} dz w(x; \hbar)$$
$$= \frac{1}{5} \log(z_1^2 z_2) - 6z_1^2 z_2 + 15z_1^2 - 2z_1 - \left( 5z_1^2 z_2 - \frac{15z_1^2}{2} + \frac{z_1^4}{4} \right) \hbar^2 + O(\hbar^4, z_1^3). \quad (4.40)$$

We note that as familiar from literature, the logarithmic term is not captured by the residue calculations and is added by hand. We express the coefficients $\Pi^{(n)}$ by the differential operator method. Since the differential operators giving $\Pi^{(n \geq 4)}$ is tedious long expression, here we provide the differential operator giving the leading correction to the classical periods as an example$^4$,

$$D_2 = \frac{1}{8} \theta_1^2 + \frac{1}{6} \theta_1 \theta_2. \quad (4.41)$$

By using the operator, we can obtain the leading correction to the quantum mirror maps $t_i^{(2)}(z; \hbar)$ and the quantum B-periods $\Pi_{d,i}^{(2)}(z; \hbar)$,

$$t_i^{(2)}(z; \hbar) = D_2 t_i(z), \quad \Pi_{d,i}^{(2)}(z; \hbar) = D_2 \Pi_{d,i}^{(0)}(z). \quad (4.42)$$

---

$^4$We provide the results of differential operators giving higher order quantum corrections in the mathematica file. The results contain the differential operators of $C^3/Z_5$ and $Y^{3,m}$ with $m = 0, 1, 2, 3$. One can find it in the source file on the arXiv.
with \( i = 1, 2 \).

To check the consistency, let us calculate the NS free energy near the large radius point. By solving the recursion relations (4.27) and (4.28) with the matrix (4.36) invertible and cancelled out, we find the NS free energy whose instanton parts \( [F_n]^{\text{inst.}} \) are given by

\[
[F_1]^{\text{inst.}} = \frac{-127Q_1^2Q_2^2}{12} - \frac{65Q_1^2Q_2}{8} + \frac{129Q_1^2}{16} + \frac{5Q_1Q_2}{6} + \frac{7Q_1}{8} - \frac{Q_2^2}{12} - \frac{Q_2}{6} + O(Q_3^2),
\]

\[
[F_2]^{\text{inst.}} = -\frac{2561Q_1^2Q_2^2}{720} - \frac{263Q_1^2Q_2}{72} + \frac{207Q_1^2}{64} + \frac{29Q_1Q_2^2}{640} + \frac{29Q_1}{640} + \frac{Q_2}{180} + \frac{Q_2}{360} + O(Q_3^2).
\]  

(4.43)

They agree with the topological vertex computations.

Now we are ready to calculate the quantum corrections to the energy spectra. The all-order Bohr–Sommerfeld quantization condition in this case is given by

\[
\text{vol}_i(E_1, E_2; \hbar) = 2\pi \hbar \left( n_i + \frac{1}{2} \right), \quad i = 1, 2,
\]  

(4.44)

where \( \text{vol}_i(E_1, E_2; \hbar) \) are the quantum corrected phase volumes. To obtain the quantum corrected spectrum, we define \( E_i \) and \( \text{vol}_i(E_1, E_2; \hbar) \) as series of \( \hbar \),

\[
E_i = \sum_{n=0}^{\infty} E_i^{(n)} \hbar^n,
\]

\[
\text{vol}_i(E_1, E_2; \hbar) = \sum_{n=0}^{\infty} \text{vol}_i^{(2n)}(E_1, E_2) \hbar^{2n}.
\]  

(4.45)

The classical Bohr–Sommerfeld volumes have to vanish in the classical limit \( \hbar = 0 \) of (4.68) at the minimum

\[
E_1^{(0)} = 5 =: E_{m_1}, \quad E_2^{(0)} = 5 =: E_{m_2},
\]  

(4.46)

which corresponds to the conifold point. By expanding (4.68) in \( \hbar \), we can obtain \( E_i^{(n)} \) as a function of \( \text{vol}_i^{(n)}(E_{m_1}, E_{m_2}) \), e.g.,

\[
E_1^{(1)} = 2\pi \left\{ \left( n_1 + \frac{1}{2} \right) \partial E_2 \text{vol}_2^{(0)} - \left( n_2 + \frac{1}{2} \right) \partial E_1 \text{vol}_1^{(0)} \right\},
\]

\[
\frac{\partial E_i \text{vol}_1^{(0)} \partial E_2 \text{vol}_2^{(0)} - \partial E_2 \text{vol}_1^{(0)} \partial E_1 \text{vol}_2^{(0)}}{\partial E_i \text{vol}_1^{(0)} \partial E_2 \text{vol}_2^{(0)} - \partial E_2 \text{vol}_1^{(0)} \partial E_1 \text{vol}_2^{(0)}}
\],

\[
E_2^{(1)} = 2\pi \left\{ \left( n_1 + \frac{1}{2} \right) \partial E_1 \text{vol}_1^{(0)} - \left( n_2 + \frac{1}{2} \right) \partial E_2 \text{vol}_2^{(0)} \right\},
\]

\[
\frac{\partial E_i \text{vol}_2^{(0)} \partial E_2 \text{vol}_1^{(0)} - \partial E_2 \text{vol}_2^{(0)} \partial E_1 \text{vol}_1^{(0)}}{\partial E_i \text{vol}_2^{(0)} \partial E_2 \text{vol}_1^{(0)} - \partial E_2 \text{vol}_2^{(0)} \partial E_1 \text{vol}_1^{(0)}}
\],

(4.47)

where we omit the arguments \( (E_{m_1}, E_{m_2}) \) of \( \text{vol}_i^{(n)} \).\footnote{We will use this expression for other models, where the arguments of \( \text{vol}_i^{(n)} \) in these models are \( (E_{m_1}, E_{m_2}, R) \).}

By comparing (4.47) with perturbative calculations (3.10), we obtain exact values of the \( E_{1,2} \)-derivatives of phase
volumes at the classical minimum,
\[
\begin{pmatrix}
\frac{\partial E_1 \text{vol}^{(0)}_1}{\partial E_1 \text{vol}^{(0)}_2} & \frac{\partial E_2 \text{vol}^{(0)}_1}{\partial E_2 \text{vol}^{(0)}_2}
\end{pmatrix} = \frac{\pi}{5} \begin{pmatrix}
-2 \left(5 - 2\sqrt{5}\right)^{1/2} & 2 \left(\sqrt{5} + 5\right)^{1/2} \\
2 \left(2\sqrt{5} + 5\right)^{1/2} & - \left(10 - 2\sqrt{5}\right)^{1/2}
\end{pmatrix}.
\] (4.48)

With the change of variables (4.38), we find
\[
\begin{pmatrix}
\frac{\partial z_1 \text{vol}^{(0)}_1(z)}{\partial z_1 \text{vol}^{(0)}_2(z)} & \frac{\partial z_2 \text{vol}^{(0)}_1(z)}{\partial z_2 \text{vol}^{(0)}_2(z)}
\end{pmatrix} = \pi \begin{pmatrix}
-10\sqrt{5} + 2\sqrt{5} & \sqrt{130 - 58\sqrt{5}} \\
-10\sqrt{5} - 2\sqrt{5} & -\sqrt{130 + 58\sqrt{5}}
\end{pmatrix}.
\] (4.49)

We check this is indeed true numerically.

The classical mirror maps near the conifold point are
\[
t_{c,1} = -2\pi \left(5 \left(2\sqrt{5} + 5\right)\right)^{1/2} z_{c,1} - \pi \left(26 - \frac{38}{\sqrt{5}}\right)^{1/2} z_{c,2} + 24\pi \left(1 - \frac{2}{\sqrt{5}}\right)^{1/2} z_{c,1} z_{c,2} + O(z_{c,i}^2),
\]
\[
t_{c,2} = -2\pi \left(50 - 10\sqrt{5}\right)^{1/2} z_{c,1} - 2\pi \left(13 + \frac{22}{\sqrt{5}}\right)^{1/2} z_{c,2} + 24\pi \left(2 + \frac{2}{\sqrt{5}}\right)^{1/2} z_{c,1} z_{c,2} + O(z_{c,i}^2),
\] (4.50)

where
\[
z_1 = \frac{1}{25} + z_{c,1}, \quad z_2 = \frac{1}{5} + z_{c,2}.
\] (4.51)

The coefficients of $z_{c,1}, z_{c,2}$ in the classical mirror map are fixed by the relation (4.22).

We can calculate the next leading order of the energy spectrum $E^{(2)}_{1,2}$ by looking at $\hbar^2$-order term of (4.44). To obtain them, we need to calculate the second derivatives of the volumes and first quantum correction which can be calculated from the formula (4.22) with (4.41). After some computations, we find
\[
E^{(2)}_1 = \frac{1}{40} \left(-2 \left(\sqrt{5} - 3\right) n_1 (n_1 + 1) + 2 \left(\sqrt{5} + 3\right) n_2 (n_2 + 1) + 7\right),
\]
\[
E^{(2)}_2 = \frac{1}{20} \left(2 \left(\sqrt{5} + 2\right) n_1^2 - 2 \left(\sqrt{5} - 2\right) n_2^2 + 4 \left(\sqrt{5} + 1\right) n_1 + 4 n_2 + 4\sqrt{5} n_1 n_2 + \sqrt{5} + 3\right).
\] (4.52)

These results agree with the perturbative computation (3.13).

**4.2.2 $Y^{3,0}$ case**

The mirror curve of $Y^{3,0}$ is
\[
e^p + z_1 z_2^2 z_3 e^{-p + 3x} + z_1 z_2^2 e^{3x} + z_2 e^{2x} + e^x + 1 = 0.
\] (4.53)

The Picard–Fuchs operators are
\[
L_1 = (\theta_1 - \theta_3)(\theta_1 - 2\theta_2) - z_1 (-2\theta_1 + \theta_2 - 1)(-2\theta_1 + \theta_2),
L_2 = (\theta_2 - \theta_3)(\theta_2 - 2\theta_1) - z_2 (-2\theta_2 + \theta_1 - 1)(-2\theta_2 + \theta_1),
L_3 = \theta_3^2 - z_3 (\theta_1 - \theta_3)(\theta_2 - \theta_3),
L_4 = \theta_3^2 - z_1 z_2 z_3 (\theta_1 - 2\theta_2)(\theta_2 - 2\theta_1).
\] (4.54)
Note that these operators are symmetric under exchange of $z_1$ and $z_2$. To give the solutions of Picard–Fuchs equation, we define following function,

$$\omega_0(\rho_i) = \sum_{l,m,n \geq 0} c(l, m, n; \rho_i) z_1^{l+\rho_1} z_2^{m+\rho_2} z_3^{n+\rho_3}$$

(4.55)

with

$$c(l, m, n; \rho_i) = \frac{1}{\Gamma(n + \rho_3 + 1) \Gamma(-n + l + \rho_1 - \rho_3 + 1) \Gamma(-n + m + \rho_2 - \rho_3 + 1)}$$

$$\times \frac{1}{\Gamma(l - 2m + \rho_1 - 2\rho_2 + 1) \Gamma(-2l + m - 2\rho_1 + \rho_2 + 1)}.$$  

(4.56)

Then, the classical mirror maps and the derivatives of the prepotential are given by

$$t_1(z) = \omega_1 = \log z_1 + 2z_1 + \frac{3}{2} z_2^2 - \frac{3}{2} z_2 z_3 + 2 z_1 z_2 z_3 + O(z_i^3),$$

$$t_2(z) = \omega_2 = t_1|_{z_1 \leftrightarrow z_2},$$

$$t_3(z) = \omega_3 = \log z_3,$$

(4.57)

and

$$\frac{\partial F_0}{\partial t_1} = \omega_{11} + \omega_{12} + \frac{1}{2} \omega_{22} + \frac{2}{3} \omega_{13} + \frac{1}{3} \omega_{23} + \frac{2\pi^2}{3},$$

$$\frac{\partial F_0}{\partial t_2} = \omega_{22} + \omega_{12} + \frac{1}{2} \omega_{11} + \frac{2}{3} \omega_{23} + \frac{1}{3} \omega_{13} + \frac{2\pi^2}{3},$$

(4.58)

where

$$\omega_i = \frac{\partial \omega_0}{\partial \rho_i} \bigg|_{\rho_{1,2,3}=0},$$

$$\omega_{ij} = \frac{\partial^2 \omega_0}{\partial \rho_i \partial \rho_j} \bigg|_{\rho_{1,2,3}=0}.$$

(4.59)

The classical B-periods $\Pi_{d,i}$ ($i = 1, 2$) are given by formula (4.4) with

$$C = \begin{bmatrix} 2 & -1 & 0 \\ -1 & 2 & 0 \end{bmatrix},$$

(4.60)

where the first 2 × 2 block is the Cartan matrix of SU(3). From the prepotential, the Bohr-Sommerfeld volumes are

$$\text{vol}^{(0)}_i(z) = \sum_{j=1}^{3} C_{ij} \frac{\partial F_0}{\partial t_j} - \frac{2\pi^2}{3}, \quad i = 1, 2,$$

(4.61)

where the complex structure moduli $z_1, z_2, z_3$ are related to the quantum systems by

$$z_1 = \frac{E_2}{E_1^2}, \quad z_2 = \frac{E_1}{E_2}, \quad z_3 = -R^6.$$

(4.62)

The classical phase volumes should vanish at the classical minimum, $z_1 = z_2 = \frac{1}{3(1+R^2)}$.

We check numerically this is indeed true for e.g. $R = 1$. 

23
Now let us consider the quantum mirror curve,

\[ \Psi(x - \imath \hbar) + z_1 z_2^2 \text{e}^{3x} x^{3\hbar} \Psi(x + \imath \hbar) + (z_1 z_2^2 \text{e}^{3x} + 2 \text{e}^{2x} + e^x + 1) \Psi(x) = 0 \]  
(4.63)

By taking the residue of \( w(x; \hbar) \), we find a quantum A-period,

\[
\Pi(z; \hbar) = -\frac{1}{3} \log(z_1 z_2^2) + \oint_{x=\infty} \text{d}x w(x; \hbar)
\]

\[
= -\frac{1}{3} \log(z_1 z_2^2) + \left( -z_2 - \frac{3z_2^2}{2} - \frac{10z_2^3}{3} + z_1 z_2^2 + 4z_1 z_2^3 + 2z_1 z_2^2 z_3 + 12z_1 z_2^3 z_3 \right)
\]

\[-\left( \frac{1}{4} z_1 z_2^2 z_3 + \frac{7}{2} z_1 z_2^3 z_3 \right) \hbar^2 + \mathcal{O}(\hbar^4, z_i^4).
\]

(4.64)

The differential operator giving the first quantum correction is

\[
\mathcal{D}_2 = \frac{1}{12} z_1 z_2 (5 z_3 + 4) \theta_1^2 + \frac{1}{12} z_1 z_2 (5 z_3 + 4) \theta_2^2 + \frac{1}{24} (-20 z_1 z_2 - 25 z_1 z_2 z_3 + 4) \theta_1 \theta_2.
\]

(4.65)

Then, we can obtain the \( \hbar^2 \) correction to the quantum mirror map and quantum B-period by acting above differential operator on the classical periods,

\[
t_i^{(2)} = \mathcal{D}_2 t_i, \quad \Pi_{i,d_i}^{(2)} = \mathcal{D}_2 \Pi_{i,d_i}^{(0)}, \quad i = 1, 2.
\]

(4.66)

We note that in this model, the \( t_3 \) depends only on mass parameter \( R \) and receives no quantum correction.

To check the consistency, let us calculate the NS free energy near the large radius point. In the similar way as in the \( \mathbb{C}^3/\mathbb{Z}_5 \) case, we can obtain the NS free energy by solving the recursion relations (4.27) and (4.28) whose instanton parts \( [F_n]^{\text{inst.}} \) are given by

\[
[F_1]^{\text{inst.}} = -\frac{1}{6} Q_1 - \frac{1}{6} Q_2 - \frac{1}{12} Q_1^2 - \frac{1}{12} Q_2^2 - \frac{1}{6} Q_1 Q_2 - \frac{1}{24} Q_1 Q_2 Q_3
\]

\[-\frac{1}{12} Q_1^2 Q_2^2 + \frac{7}{8} Q_1^2 Q_2 Q_3 + \frac{7}{8} Q_1 Q_2^2 Q_3 + \frac{5}{6} Q_1 Q_2 Q_3^2 - \frac{1}{48} Q_1^2 Q_2^2 Q_3^2 + \mathcal{O}(Q_i^3),
\]

\[
[F_2]^{\text{inst.}} = \frac{1}{360} Q_1 + \frac{1}{180} Q_2 + \frac{1}{180} Q_1^2 + \frac{1}{180} Q_2^2 + \frac{1}{360} Q_1 Q_2 + \frac{1}{5760} Q_1 Q_2 Q_3
\]

\[+ \frac{1}{180} Q_1^2 Q_2^2 + \frac{29}{640} Q_1^2 Q_2 Q_3 + \frac{29}{640} Q_1 Q_2^2 Q_3 + \frac{37}{1440} Q_1^2 Q_2^2 Q_3 + \frac{7}{2880} Q_1^2 Q_2^2 Q_3^2 + \mathcal{O}(Q_i^3).
\]

(4.67)

They agree with the topological vertex computations. Accidentally, it turns out that the derivatives with mass parameter \( \partial_{\theta_i} F_1 \) also satisfies a similar equation although it does not formally appear in (4.27) for this model.

Now we are ready to calculate the quantum corrections to the energy spectra. The all-order Bohr–Sommerfeld quantization conditions are given by

\[
\text{vol}_i(E_1, E_2, R; \hbar) = 2\pi \hbar \left( n_i + \frac{1}{2} \right), \quad i = 1, 2,
\]

(4.68)
where \( \text{vol}_i(E_1, E_2, R; \hbar) \) are the quantum corrected phase volume. To obtain the quantum corrected spectrum, we define \( E_i \) and \( \text{vol}_i(E_1, E_2, R; \hbar) \) as series of \( \hbar \),

\[
E_i = \sum_{n=0}^{\infty} E_i^{(n)} \hbar^n, \tag{4.69}
\]

\[
\text{vol}_i(E_1, E_2, R; \hbar) = \sum_{n=0}^{\infty} \text{vol}_i^{(n)}(E_1, E_2, R) \hbar^{2n}. \]

The classical Bohr–Sommerfeld volumes have to vanish in the classical limit \( \hbar = 0 \) of (4.68) at the minimum

\[
E_i^{(0)} = 3(1 + R^2) =: E_m, \tag{4.70}
\]

which corresponds to the conifold point. In the following, we demonstrate the computation for \( R = 1 \). The leading corrections to the spectra are given by (4.47). By comparing them with direct perturbative calculations (3.14), we obtain the exact value of \( E_{1,2} \)-derivatives of the volumes,

\[
\left( \frac{\partial E_1 \text{vol}_1^{(0)}}{\partial E_2 \text{vol}_1^{(0)}}, \frac{\partial E_2 \text{vol}_1^{(0)}}{\partial E_1 \text{vol}_1^{(0)}} \right) = \pi \left( \left( \frac{2}{15} (\sqrt{5} + 3) \right)^{1/2} - \left( \frac{2}{15} (3 - \sqrt{5}) \right)^{1/2} \right). \tag{4.71}
\]

With the changes of variables, we find

\[
\left( \frac{\partial z_1 \text{vol}_1^{(0)}(z)}{\partial z_2 \text{vol}_1^{(0)}(z)}, \frac{\partial z_2 \text{vol}_1^{(0)}(z)}{\partial z_3 \text{vol}_1^{(0)}(z)} \right) = -\frac{8\sqrt{3}\pi}{3 - \sqrt{5}} \begin{pmatrix} 1 - \sqrt{5} & -5 + 2\sqrt{5} \\ -5 + 2\sqrt{5} & 1 - \sqrt{5} \end{pmatrix}, \tag{4.72}
\]

They agree with the direct computation numerically.

To obtain the derivatives of the volumes, we use the classical periods near the conifold point,

\[
t_{c,1} = -\frac{4\pi(5 + 9\sqrt{5})}{5\sqrt{3}} z_{c,1} + \frac{4\pi(5 - 9\sqrt{5})}{5\sqrt{3}} z_{c,2} + z_{c,3} + \frac{1312\pi}{25\sqrt{15}} z_{c,1} z_{c,2} + \frac{8\pi(125 + 117\sqrt{5})}{1125\sqrt{3}} z_{c,1} z_{c,3} - \frac{8\pi(125 - 117\sqrt{5})}{1125\sqrt{3}} z_{c,2} z_{c,3} - \frac{4544\pi}{1125\sqrt{15}} z_{c,1} z_{c,2} z_{c,3} + O(z_{c,i}^2),
\]

\[
t_{c,2} = t_{c,1}|_{z_{c,1} \rightarrow z_{c,2}},
\]

\[
t_{c,3} = \log(-1 + z_{c,3}), \tag{4.73}
\]

where

\[
z_1 = \frac{1}{6} + z_{c,1}, \quad z_2 = \frac{1}{6} + z_{c,2}, \quad z_3 = -1 + z_{c,3}. \tag{4.74}
\]

The coefficients of \( z_{c,1} \) and \( z_{c,2} \) are fixed by the relation (4.22).

From them, we can obtain the next leading order of the quantum corrections to the energy spectra by looking at \( \hbar^2 \)-order of (4.68). After some computations, we have

\[
E_1^{(2)} = \frac{1}{360} \left( 6(19 + 5\sqrt{5})n_1 + 6(19 - 5\sqrt{5})n_2 + 6(13 + 5\sqrt{5})n_1^2 + 6(13 - 5\sqrt{5})n_1^2 + 72n_1 n_2 + 101 \right),
\]

\[
E_2^{(2)} = E_1^{(2)}|_{n_1 \leftrightarrow n_2}. \tag{4.75}
\]
They agree with the perturbative computation given in (3.14).

4.2.3 $Y^{3,1}$ case

In this example, we sometimes use some of the notations and definitions given in 4.2.2. The mirror curve of $Y^{3,1}$ is,

$$e^p + z_3 e^{2x-p} + z_1 z_2^2 e^{3x} + z_2 e^{2x} + e^x + 1 = 0. \quad (4.76)$$

The Picard–Fuchs operators are

$$L_1 = \theta_1(\theta_1 - 2\theta_2 - 2\theta_3) - z_1(-2\theta_1 + \theta_2 - 1)(-2\theta_1 + \theta_2),$$
$$L_2 = \theta_2(-2\theta_1 + \theta_2) - z_2(\theta_1 - 2\theta_2 - 2\theta_3 - 1)(\theta_1 - 2\theta_2 - 2\theta_3),$$
$$L_3 = \theta_3^2 - z_3(\theta_1 - 2\theta_2 - 2\theta_3 - 1)(\theta_1 - 2\theta_2 - 2\theta_3),$$
$$L_4 = \theta_1\theta_2\theta_3^2 - z_1 z_2 z_3(-2\theta_1 + \theta_2)(\theta_1 - 2\theta_2 - 2\theta_3 - 2)(\theta_1 - 2\theta_2 - 2\theta_3 - 1)(\theta_1 - 2\theta_2 - 2\theta_3). \quad (4.77)$$

Then, the classical mirror maps and the derivatives of the prepotential are given by

$$t_1(z) = \omega_1 = \log z_1 + 2z_1 - z_2 - z_3 + 3z_1^2 - \frac{3z_1^2}{2} - \frac{3z_2^2}{2} - 6z_2 z_3 - 2z_1 z_2 + 6z_1^2 z_3 + z_1 z_2^2 - 30z_2^2 z_3 - 30z_2 z_3^2 - 315z_2^2 z_3^2 + 12z_1 z_2^2 z_3 + 90z_1 z_2^2 z_3^2 + O(z_i^3),$$
$$t_2(z) = \omega_2 = \log z_2 + 2z_2 - z_1 + 2z_2 + 2z_3 - \frac{3z_1^2}{2} + 3z_2^2 + 3z_3^2 + 12z_2 z_3 + z_1^2 z_2 - 3z_1^2 z_3 - 2z_1 z_2^2 + 60z_2^2 z_3 + 60z_2 z_3^2 + 630z_2^2 z_3^2 - 24z_1 z_2^2 z_3 - 180z_1 z_2^2 z_3^2 + O(z_i^3),$$
$$t_3(z) = \omega_3 = \log z_3 + \frac{1}{3} \{ 2 (t_1(z) - \log z_1) + 4 (t_2(z) - \log z_2) \}, \quad (4.78)$$

and

$$\frac{\partial F_0}{\partial t_1} = \frac{1}{9} \left( 4 \omega_{11} - 2 \omega_{12} - 2 \omega_{22} + 6 \omega_{13} + 3 \omega_{23} \right),$$
$$\frac{\partial F_0}{\partial t_2} = \frac{1}{9} \left( - \omega_{11} - 4 \omega_{12} - 4 \omega_{22} + 3 \omega_{13} + 6 \omega_{23} \right), \quad (4.79)$$
$$\frac{\partial F_0}{\partial t_3} = \frac{1}{3} \left( \omega_{11} + \omega_{22} + \omega_{12} \right),$$

where $\omega_i$ and $\omega_{ij}$ are defined in (4.59), and $\omega_0$ are defined in (4.55) with the coefficient $c(l, m, n; \rho_i)$

$$c(l, m, n; \rho_i) = \frac{\Gamma (\rho_1 + 1) \Gamma (\rho_2 + 1) \Gamma (\rho_3 + 1)}{\Gamma (l + \rho_1 + 1) \Gamma (m + \rho_2 + 1) \Gamma (n + \rho_3 + 1)} \times \frac{\Gamma (-2 \rho_1 + \rho_2 + 1) \Gamma (\rho_1 - 2 \rho_2 - 2 \rho_3 + 1)}{\Gamma (l + \rho_1 - 2 (m + \rho_2) - 2 (n + \rho_3) + 1) \Gamma (m - 2 (l + \rho_1) + \rho_2 + 1)}. \quad (4.80)$$

The classical B-periods $\Pi_{d,i}$ ($i = 1, 2$) are given by formula (4.4) with the non-invertible matrix

$$C = \begin{bmatrix} 2 & -1 & 0 \\ -1 & 2 & 2 \end{bmatrix}. \quad (4.81)$$

26
From the prepotential, the Bohr-Sommerfeld volumes are
\[ \text{vol}_i^{(0)}(z) = \sum_{j=1}^{3} C_{ij} \frac{\partial F_0}{\partial t_j} - \frac{2\pi^2}{3}, \quad i = 1, 2. \] (4.82)

The complex structure moduli \( z_1, z_2, z_3 \) are related to the dimer system as following,
\[ z_1 = \frac{E_2}{E_1^2}, \quad z_2 = \frac{E_1}{E_2}, \quad z_3 = \frac{R^6}{E_2^2}. \] (4.83)

From (3.18), the classical phase volumes vanish at
\[ z_1 = \frac{r^9 + 2}{(r^9 - 4)^2}, \quad z_2 = -\frac{r^{18}(r^4 - 4)}{(r^9 + 2)^2}, \quad z_3 = -\frac{(r^9 - 1)^3}{(r^9 + 2)^2}. \] (4.84)

where we use the polynomial relation (3.17) to eliminate \( R \) and write the expressions purely in terms of \( r \). As an example, for \( r = 2^{-1/9} \), we check numerically this is indeed true for (4.82).

Now let us consider the quantum mirror curve,
\[ \Psi(x - i\hbar) + e^{i\hbar z_3 e^{2x}} \Psi(x + i\hbar) + \left( z_1 z_2^2 e^{3x} + z_2 e^{2x} + e^x + 1 \right) \Psi(x) = 0 \] (4.85)

By taking the residue of \( w(x; \hbar) \), we find a quantum A-period,
\[ \Pi(z; \hbar) = -\frac{1}{3} \log(z_1 z_2^2) + \oint_{x=\infty} dw(x; \hbar) \]
\[ = -\frac{1}{3} \log(z_1 z_2^2) + \left( -z_2 - z_3 - 6z_2 z_3 - \frac{3z_2^2}{2} - \frac{3z_3^2}{2} - z_1 z_2 - 30z_2 z_3 - 30z_2 z_3^2 \right) \]
\[ - 315z_2^2 z_3^3 + 90z_1 z_2^2 z_3^2 + 12z_1 z_2 z_3^2 \]
\[ - (-z_2 z_3 - 10z_2 z_3^2 - 10z_2^2 z_3 - 210z_2 z_3^2 + 6z_1 z_2 z_3 + 90z_1 z_2 z_3^2) \hbar^2 + O(\hbar^4, z_1^3). \] (4.86)

The quantum corrections can be expressed as the classical part acted by the differential operator,
\[ D_2 = -\frac{z_1}{2} \theta_1 + \left( \frac{1}{12} - z_1 \right) \theta_1^2 + \frac{z_1}{4} \theta_2 + \left( \frac{1}{6} - \frac{z_1}{4} \right) \theta_2^2 + \left( -\frac{1}{12} + z_1 - \frac{1}{12z_1 z_2} \right) \theta_1 \theta_2. \] (4.87)

Then, we can obtain the quantum mirror map and quantum B-period by acting the differential operator on the classical periods,
\[ t_i^{(2)} = D_2 t_i, \quad (i = 1, 2, 3), \quad \Pi_{d,k}^{(2)} = D_2 \Pi_{d,k}^{(0)}, \quad (k = 1, 2). \] (4.88)

The NS free energy near the large radius point can be calculated from the general formulae (4.27) and (4.28) whose instanton parts \([F_n]^{\text{inst.}} \) are given by
\[ [F_1]^{\text{inst.}} = \frac{Q_1}{6} + \frac{Q_2}{6} + \frac{Q_3}{6} + \frac{Q_1 Q_2}{6} + \frac{Q_1 Q_3}{6} + \frac{7Q_2 Q_3}{3} + \frac{5Q_1 Q_2 Q_3}{2} + O(Q_1^2), \]
\[ [F_2]^{\text{inst.}} = \frac{Q_1}{360} + \frac{Q_2}{360} + \frac{Q_3}{360} + \frac{Q_1 Q_2}{360} + \frac{Q_1 Q_3}{360} + \frac{59Q_2 Q_3}{180} - \frac{13Q_1 Q_2 Q_3}{40} + O(Q_1^2). \] (4.89)
They agree with the topological vertex computations.

Now we are ready to calculate the quantum corrections to the energy spectra. The Bohr–Sommerfeld quantization condition is given by (4.68) with the quantum corrected spectra and volumes defined in (4.69). The classical Bohr–Sommerfeld volumes have to vanish in the classical limit \( \hbar = 0 \) of (4.68) at the minimum

\[
E_1^{(0)} = \frac{4 - r^9}{r^6} =: E_{m_1}, \quad E_2^{(0)} = \frac{2 + r^9}{r^{12}} =: E_{m_2}
\]

which corresponds to the conifold point. In the followings, we do the computations for a particular case \( r = 2^{-1/9} \). The leading corrections to the spectra are given by (4.47). By comparing with the perturbative computation (3.18), we find the exact values of \( E_{1,2} \)-derivatives of the volumes at the conifold point,

\[
\begin{pmatrix}
\partial_{E_1} \text{vol}_1^{(0)} & \partial_{E_2} \text{vol}_1^{(0)} \\
\partial_{E_1} \text{vol}_2^{(0)} & \partial_{E_2} \text{vol}_2^{(0)}
\end{pmatrix} = \pi \begin{pmatrix}
-\frac{2\sqrt{2}}{2^{2/3} (166\sqrt{2}+245)^{1/6}} & \frac{1}{2^{11/12} (\sqrt{3}+1) (2^{1/4} (166\sqrt{2}+245)^{1/6} - (3\sqrt{2}-52)^{1/2})} \\
-\frac{2^{1/3}}{4^{1/2} (2^{1/4} (166\sqrt{2}+245)^{1/6} - (3\sqrt{2}-52)^{1/2})} & \frac{2^{23/6} (7\sqrt{2}+8)^{1/2} (166\sqrt{2}+245)^{1/6} - 2(31-34\sqrt{2})^{1/2}}{2^{11/12} (\sqrt{3}+1) (2^{1/4} (166\sqrt{2}+245)^{1/6} - (3\sqrt{2}-52)^{1/2})}
\end{pmatrix},
\]

which agree with the numerical computation.

In this case we do not calculate the classical mirror map around the conifold point, but when one wants to calculate higher corrections to the energy spectra as in the case of \( Y^{3,0} \), the classical mirror map is needed to obtain the higher order quantum corrections to the (derivatives) of the volumes via the formulae (4.18), (4.22).

### 4.2.4 \( Y^{3,2} \) case

In this example, we sometimes also use some of the notations and definitions given in section 4.2.2. The mirror curve of \( Y^{3,2} \) is

\[
e^x + z_3 e^{x-p} + z_1 z_2^2 e^{3x} + z_2 e^{2x} + e^x + 1 = 0.
\]

(4.92)

The Picard–Fuchs operators are

\[
\begin{align*}
\mathcal{L}_1 &= \theta_1 (\theta_1 - 2\theta_2 - \theta_3) - z_1(-2\theta_1 + \theta_2 - 1)(-2\theta_1 + \theta_2), \\
\mathcal{L}_2 &= (\theta_2 - \theta_3)(-2\theta_1 + \theta_2) - z_2(\theta_1 - 2\theta_2 - \theta_3 - 1)(\theta_1 - 2\theta_2 - \theta_3), \\
\mathcal{L}_3 &= \theta_2^2 - z_3(\theta_1 - 2\theta_2 - \theta_3)(\theta_2 - \theta_3), \\
\mathcal{L}_4 &= \theta_1 \theta_3^2 - z_1 z_2^2(\theta_1 - 2\theta_2 - \theta_3 - 1)(\theta_1 - 2\theta_2 - \theta_3)(-2\theta_1 + \theta_2).
\end{align*}
\]

(4.93)

The classical mirror maps and the derivatives of the prepotential are given by

\[
\begin{align*}
t_1(z) &= \omega_1 = \log z_1 + 2z_1 - z_2 - 3z_1^2 - \frac{3z_1^2}{2} + 2z_2 z_3 - 2z_1^2 z_2 + z_1 z_2^2 + 12z_2^2 z_3 \\
&\quad - 15z_2 z_3^2 - 6z_1 z_2^2 z_3 + 6z_1 z_2 z_3^2 + \mathcal{O}(z_1^3), \\
t_2(z) &= \omega_2 = \log z_2 - z_1 + 2z_2 - \frac{3z_1^2}{2} + 3z_2^2 + 4z_2 z_3 + z_1^2 z_2 - 2z_1 z_2^2 - 24z_2^2 z_3 \\
&\quad + 30z_2 z_3^2 + 12z_1 z_2^2 z_3 - 12z_1 z_2 z_3^2 + \mathcal{O}(z_1^3), \\
t_3(z) &= \omega_3 = \log z_3 + \frac{1}{3} \left\{ (t_1(z) - \log z_1) + 2 (t_2(z) - \log z_2) \right\},
\end{align*}
\]

(4.94)
and
\[ \frac{\partial F_0}{\partial t_1} = \frac{1}{18} (16\omega_{11} + 10\omega_{12} + \omega_{22} + 12\omega_{13} + 6\omega_{23}), \]
\[ \frac{\partial F_0}{\partial t_2} = \frac{1}{18} (5\omega_{11} + 2\omega_{12} + 2\omega_{22} + 6\omega_{13} + 12\omega_{23}), \]
\[ \frac{\partial F_0}{\partial t_3} = \frac{1}{3} (\omega_{11} + \omega_{22} + \omega_{12}) \]

where \( \omega_i \) and \( \omega_{ij} \) are defined in (4.59) with the coefficient \( c(l, m, n; \rho_i) \) given by

\[ c(l, m, n; \rho_i) = \frac{\Gamma (\rho_1 + 1) \Gamma (\rho_3 + 1)^2 \Gamma (\rho_2 - \rho_3 + 1)}{\Gamma (l + \rho_1 + 1) \Gamma (n + \rho_3 + 1)^2 \Gamma (m - n + \rho_2 - \rho_3 + 1)} \times \frac{\Gamma (-2\rho_1 + \rho_2 + 1) \Gamma (\rho_1 - 2\rho_2 - \rho_3 + 1)}{\Gamma (m - 2(l + \rho_1) + \rho_2 + 1) \Gamma (l - n + \rho_1 - 2(m + \rho_2) - \rho_3 + 1)}. \]

The classical B-periods \( \Pi_{d,i} \) \( (i = 1, 2) \) are given by (4.4) with the non-invertible matrix \( C_{ij} \),

\[ C = \begin{bmatrix} 2 & -1 & 0 \\ -1 & 2 & 1 \end{bmatrix}. \]

From the prepotential, the Bohr-Sommerfeld volumes are

\[ \text{vol}^{(0)}_i(z) = \sum_{i=1}^{3} C_{ij} \frac{\partial F_0}{\partial t_j} - \frac{2\pi^2}{3}, \quad i = 1, 2. \]

The complex structure moduli \( z_1, z_2, z_3 \) are related to the dimer model by

\[ z_1 = \frac{E_2}{E_1}, \quad z_2 = \frac{E_1}{E_2}, \quad z_3 = -\frac{R^6}{E_2}. \]

From (3.21), the classical phase volumes vanish at

\[ z_1 = \frac{r^{9/2} - 3r^{9/4} + 5}{(2r^{9/4} - 5)^2}, \quad z_2 = \frac{5r^{9/4} - 2r^{9/2}}{(r^{9/2} - 3r^{9/4} + 5)^2}, \quad z_3 = \frac{(r^{9/4} - 1)^3}{r^{9/4}(r^{9/2} - 3r^{9/4} + 5)}, \]

where we use the polynomial relation (3.20) to eliminate \( R \). We check numerically that the B-periods vanish at this point for e.g. \( r = 2^{-4/9} \).

Now let us consider the quantum mirror curve,

\[ \Psi(x - i\hbar) + e^{i\frac{\pi}{4}}z_3e^{x}\Psi(x + i\hbar) + (z_1z_2^2e^{3x} + z_2e^{2x} + e^{x} + 1) \Psi(x) = 0 \]

By taking the residue of \( w(x; \hbar) \), we find a quantum A-period,

\[ \Pi(z; \hbar) = -\frac{1}{3} \log(z_1z_2^2) + \oint_{x=\infty} dxw(x; \hbar) \]
\[ = -\frac{1}{3} \log(z_1z_2^2) + \left( -z_2 - \frac{3z_2^2}{2} + 2z_2z_3 + z_1z_2^2 + 12z_2^2z_3 - 15z_2^2z_3^2 - 6z_1z_2^2z_3 + 6z_1z_2^2z_3^2 \right) \]
\[ - \left( \frac{z_2z_3}{4} + \frac{7z_2^2z_3}{2} - \frac{15z_2^2z_3^2}{2} - \frac{11}{4} z_1z_2^2z_3 + 5z_1z_2^2z_3^2 \right) \hbar^2 + O(\hbar^4, z_i^3). \]
The differential operator in this case is relatively long expression,

\[ D_2 = -\frac{1}{24z_2(1-z_3)} \left\{ 2(4 - 5z_3 + 12z_1z_2 - 13z_1z_2z_3)\theta_1 \right. \\
+ \frac{1}{z_1} \left( -4 + 16z_1 + 5z_3 - 4z_1z_2 - 20z_1z_3 + 48z_1^2z_2 + 3z_1z_2z_3 - 52z_1^2z_2z_3 \right)\theta_1^2 \\
+ (-4 + 5z_3 - 12z_1z_2 + 13z_1z_2z_3)\theta_2 \\
+ (4 + 8z_2 - 5z_3 + 12z_1z_2 - 12z_2z_3 - 13z_1z_2z_3)\theta_2^2 \\
+ \frac{1}{z_1} \left( 12 - 16z_1 - 15z_3 + 4z_1z_2 + 20z_1z_3 - 48z_1^2z_2 + 52z_1^2z_2z_3 \right) \theta_1 \theta_2 \left\} \right. \\
\left. (4.103) \right. \]

Then, we can obtain the quantum mirror maps and quantum B-periods by acting above operator on the classical periods, as in (4.88).

We do not write down the NS free energy in this case since the computation process is completely the same as the case of \( Y^{3,1} \), but one can show that the NS free energy calculated from the differential operators agree with the topological vertex computations.

Now we are ready to calculate the quantum corrections to the energy spectra. The Bohr–Sommerfeld quantization condition is given by (4.68) with the quantum corrected spectra and volumes defined in (4.69). The classical Bohr–Sommerfeld volumes have to vanish in the classical limit \( \hbar = 0 \) of (4.68) at the classical minimum

\[ E_{1}^{(0)} = 5 - \frac{r^{3/4}}{r^{3/4}} =: E_{m1}^{(0)}, \quad E_{2}^{(0)} = \frac{5}{r^{3/2}} - 3r^{3/4} + r^{3} =: E_{m2}^{(0)} \] (4.104)

which corresponds to the conifold point. For simplicity, we do the computation for \( r = 2^{-4/9} \). The leading corrections to the spectra are given by (4.47). By comparing with the perturbative computation, we find the exact values of \( E_{1,2}^{(0)} \)-derivatives of the volumes at the conifold point,

\[ \begin{pmatrix} \partial_{E_1} \text{vol}_1^{(0)} \\ \partial_{E_1} \text{vol}_2^{(0)} \end{pmatrix} = \frac{\pi}{3} \begin{pmatrix} 2^{11/3} & 2^{7/3} \\ \sqrt[3]{7} & -2^{2/3} \end{pmatrix} \]

(4.105)

which are consistent with the numerical computation.

Similar to the \( Y^{3,1} \) case, we do not calculate the classical mirror map around the conifold point, but when one wants to calculate higher corrections to the energy spectra as in the case of \( Y^{3,0} \), the classical mirror map is needed to obtain the higher order quantum corrections to the (derivatives) of the volumes via the formulae (4.18) (4.22).
4.2.5 \( Y^{3,3} \) case

As the final example, we consider the \( Y^{3,3} \) model. We again sometimes use some of the notation and definition given in section 4.2.2. The mirror curve of \( Y^{3,3} \) is

\[
e^p + e^{-p} + e^{3x} + \frac{e^{2x}}{z_1^{1/3} z_2^{2/3} z_3^{1/6}} + \frac{e^x}{z_1^{2/3} z_2^{1/3} z_3^{1/3}} + \frac{1}{z_3^{1/2}} = 0. \tag{4.106}
\]

The Picard–Fuchs operators are

\[
\mathcal{L}_1 = \theta_1 (\theta_1 - 2\theta_2 - 2\theta_3) + 4\theta_2\theta_3 - z_1 (2\theta_1 - \theta_2 + 1) (2\theta_1 - \theta_2), \\
\mathcal{L}_2 = \theta_2 (\theta_2 - 2\theta_1) + z_2 (2\theta_2 - \theta_1 + 1) (2\theta_2 - \theta_1), \\
\mathcal{L}_3 = \theta_3^2 + z_3 (2\theta_3 - \theta_1 + 1) (2\theta_3 - \theta_1), \\
\mathcal{L}_4 = \theta_3^2 z_3 (\theta_1 - 2\theta_2) (\theta_1 - 3\theta_3) (2\theta_1 - \theta_2).
\tag{4.107}
\]

The solutions provide the mirror maps and the derivatives of prepotential,

\[
t_1(z) = \omega_1 = \log z_1 + 2z_1 - z_2 - z_3 + 3z_1^2 - 3z_2^2 - 3z_3^2 - 2z_1^2z_2 + 6z_1^2z_3 + z_1z_2^2 - 4z_1^2z_2z_3 + O(z_3^3),
\]

\[
t_2(z) = \omega_2 = \log z_2 - z_1 + 2z_2 - 3z_1^2 - 3z_2^2 + z_1^2z_2 - 3z_1^2z_3 - 2z_1z_2^2 + 2z_1^2z_2z_3 + O(z_3^3),
\]

\[
t_3(z) = \omega_3 = \log (z_3) + 2z_3 + 3z_3^2 + O(z_3^3),
\tag{4.108}
\]

and

\[
\frac{\partial F_0}{\partial t_1} = \frac{2}{3} \omega_{11} + \frac{2}{3} \omega_{12} + \frac{2}{3} \omega_{13} + \frac{2}{3} \omega_{22} + \frac{1}{3} \omega_{23} + \frac{2\pi^2}{3}, \\
\frac{\partial F_0}{\partial t_2} = \frac{1}{3} \omega_{11} + \frac{4}{3} \omega_{12} + \frac{1}{3} \omega_{13} + \frac{4}{3} \omega_{22} + \frac{2}{3} \omega_{23} + \frac{2\pi^2}{3},
\tag{4.109}
\]

where \( \omega_i \) and \( \omega_{ij} \) are given in (4.59) with the coefficient \( c(l, m, n; \rho_i) \),

\[
c(l, m, n; \rho_i) = \frac{1}{\Gamma(1 + l - 2m + \rho_1 - 2\rho_2) \Gamma(1 - 2l + m - 2\rho_1 + \rho_2) \Gamma(1 + l - 2n + \rho_1 - 2\rho_3)} \\
\times \frac{1}{\Gamma(1 + m + \rho_2) \Gamma(1 + n + \rho_3) \Gamma(1 + m + \rho_2) \Gamma(1 + n + \rho_3) \Gamma(1 + l - 2n + \rho_1 - 2\rho_3)}.
\tag{4.110}
\]

For the third mirror map \( t_3 \), we can calculate the summation exactly,

\[
t_3(z) = \log z_3 - 2 \log \left( \frac{1 - \sqrt{1 - 4z_3}}{2} \right). \tag{4.111}
\]

The classical B-periods are completely the same form as the one of \( Y^{3,0} \) since the matrices \( C_{ij} \) of \( Y^{3,0} \) and \( Y^{3,3} \) are the same. From the prepotential, the Bohr-Sommerfeld volumes are

\[
\text{vol}_i^{(0)}(z) = \sum_{j=1}^3 C_{ij} \frac{\partial F_0}{\partial t_j} - \frac{2\pi^2}{3}, \quad i = 1, 2.
\tag{4.112}
\]
where the complex structure moduli \( z_1, z_2, z_3 \) are related to the dimer model by
\[
 z_1 = \frac{(1 + R^6)E_1}{E_2^5}, \quad z_2 = \frac{E_2}{E_1^2}, \quad z_3 = \frac{R^6}{(1 + R^6)^2}.
\] (4.113)

The Bohr-Sommerfeld volumes should vanish at the conifold point,
\[
 z_1 = \frac{(1 + R^2)E_1}{3(1 + R^2 + R^4)^2}, \quad z_2 = \frac{1 + R^2 + R^4}{3(1 + R^2)^2}, \quad z_3 = -\frac{R^6}{(1 + R^6)^2}.
\] (4.114)

We check that the volumes vanish numerically for e.g. \( R = 1 \).

Now let us quantize the mirror curve. Correspondingly, the classical mirror curve is replaced by the difference equation,
\[
 \Psi(x + i\hbar) + \Psi(x - i\hbar) + \left( e^{3x} + \frac{e^{2x}}{z_1^{1/3} z_2^{2/3} z_3^{1/6}} + \frac{e^x}{z_1^{2/3} z_2^{1/3} z_3^{1/3}} + \frac{1}{z_3^{1/2}} \right) \Psi(x) = 0,
\] (4.115)

By taking the residue of \( w(x; \hbar) \), we find a quantum A-period,
\[
 \Pi(z; \hbar) = -\frac{1}{3} \log(z_1^2 z_2 z_3) + \oint_{x=\infty} dx w(x; \hbar)
 = -\frac{1}{3} \log(z_1^2 z_2 z_3) + \left( 2 z_1^2 z_2 z_3 + z_1^2 z_2 - 3 z_1^2 z_3 - \frac{3 z_1^2}{2} - z_1 \right)
 - (z_1^2 z_3 - z_1^2 z_2 z_3) \hbar^2 + O(\hbar^4, z_3^3).
\] (4.116)

The \( \hbar \)-corrections can be expressed as the classical part acted by the differential operator,
\[
 D_2 = \frac{1}{12(-1 + 2z_2)} \left( \frac{z_1 (15 z_2^2 - 12 z_2 + 4) - z_2 \theta_1}{2} + \frac{z_1 (9 z_2^2 - 24 z_2 + 8) + 5 z_2 - 2 \theta_1^2}{2} \right)
 + (z_1 (-15 z_2^2 + 6 z_2 - 1) + z_2) \theta_2 - \frac{z_1 (36 z_2^2 - 15 z_2 + 2) - 4 z_2 + 1 \theta_2^2}{2}.
\] (4.117)

Then, we can obtain the quantum B-period by acting above differential operator on the classical B-period.

We do not write down the NS free energy in this case since the computation process is completely the same as the case of \( Y^{3,0} \), but one can show that the NS free energy calculated from the differential operators agrees with the topological vertex computations.

Now we are ready to calculate the quantum corrections to the energy spectra. In the following computation, we consider the particular case of \( R = 1 \) for simplicity. The Bohr-Sommerfeld quantization condition is given by (4.68) with the quantum corrected spectra and volumes defined in (4.69).

The classical Bohr-Sommerfeld volumes have to vanish in the classical limit \( \hbar = 0 \) of (4.68) at the classical minimum
\[
 E_1^{(0)} = 6 =: E_{m_1}, \quad E_2^{(0)} = 9 =: E_{m_2},
\] (4.118)
which corresponds to the conifold point. The leading corrections to the spectra are
given by (4.47). By comparing them with direct perturbative calculations (3.15), we
find the exact values of $E_1, 2$-derivatives of the volumes at conifold point,
\begin{equation}
\begin{pmatrix}
\partial E_1 \vol_1^{(0)} & \partial E_2 \vol_1^{(0)} \\
\partial E_1 \vol_2^{(0)} & \partial E_2 \vol_2^{(0)}
\end{pmatrix} = \begin{pmatrix}
\frac{\sqrt{3}}{\sqrt{3}} & -\frac{1}{\sqrt{3}} \\
-\frac{1}{\sqrt{3}} & \frac{\sqrt{3}}{\sqrt{3}}
\end{pmatrix}.
\end{equation}

With the change of variables to complex structure moduli, we find
\begin{equation}
\begin{pmatrix}
\partial z_1 \vol_1^{(0)}(z) & \partial z_2 \vol_1^{(0)}(z) \\
\partial z_1 \vol_2^{(0)}(z) & \partial z_2 \vol_2^{(0)}(z)
\end{pmatrix} = \sqrt{3} \pi \begin{pmatrix}
-9 & 4 \\
0 & -12
\end{pmatrix}.
\end{equation}

The classical A-periods near the conifold point are
\begin{equation}
t_{e,1} = -3 \pi \sqrt{3} z_{c,1} - \frac{68 \pi z_{c,2}}{3 \sqrt{3}} - \frac{20 \pi z_{c,3}}{9 \sqrt{3}} + \frac{131 \pi z_{c,2} z_{c,2}}{2 \sqrt{3}} + \frac{7 \pi z_{c,1} z_{c,3}}{2 \sqrt{3}} + \frac{400 \pi z_{c,2} z_{c,3}}{27 \sqrt{3}} - \frac{1657 \pi z_{c,1} z_{c,2} z_{c,3}}{36 \sqrt{3}}
+ O(z_{c,1}^2),
\end{equation}
\begin{equation}
t_{e,2} = -6 \pi \sqrt{3} z_{c,1} - \frac{28 \pi z_{c,2}}{3 \sqrt{3}} - \frac{28 \pi z_{c,3}}{9 \sqrt{3}} + \frac{19 \pi z_{c,2} z_{c,2}}{2 \sqrt{3}} + \frac{23 \pi z_{c,1} z_{c,3}}{2 \sqrt{3}} + \frac{80 \pi z_{c,2} z_{c,3}}{27 \sqrt{3}} + \frac{7 \pi z_{c,1} z_{c,2} z_{c,3}}{36 \sqrt{3}}
+ O(z_{c,1}^2),
\end{equation}
\begin{equation}
t_{e,3} = -2 \log(1 - 2 \sqrt{-z_{c,3}}) + \log(1 + 4 z_{c,3}),
\end{equation}
where
\begin{equation}
z_1 = \frac{1}{6} + z_{c,1}, \quad z_2 = \frac{1}{6} + z_{c,2}, \quad z_3 = -1 + z_{c,3}.
\end{equation}
The coefficients of $z_{c,1}$ and $z_{c,2}$ are fixed by the relation (4.22).

Repeating the computation for $\hbar^2$-order, we find,
\begin{equation}
E_1^{(2)} = \frac{1}{36} (9 n_1 (n_1 + 1) + 3 n_2 (n_2 + 1) + 8),
\end{equation}
\begin{equation}
E_2^{(2)} = \frac{1}{2} n_1 + \frac{n_2^2}{2} + \frac{3 n_2}{2} + n_1 n_2 + \frac{2}{3}.
\end{equation}
These results agree with (3.15) with $R = 1$.

5 Discussions

In this paper, we studied the analytic connections between genus-2 mirror curves and
$Y^{p,q}$ cluster integrable systems, which are generalizations of affine $A$-type relativistic
Toda systems. It is interesting to consider the more higher genus mirror curves and
the application to other types affine Toda systems.

In the topic of the differential operator method, there are still interesting issues to
be clarify. For example, it would be interesting to consider the genus-1 mirror curves
for local $E_n$ del Pezzo surfaces, where the global symmetries are $E_n$ groups. Such
curves are considered in [52, 6] with some mass parameters turning off. With all mass
parameters turning on, the Calabi-Yau threefolds are non-toric, it is interesting to study the differential operator approach for these cases [53].

Also, in [54], the authors pointed out that the quantum A-periods of $D_5$ del Pezzo geometry can be expressed as $D_5$ Weyl characters. The quantum mirror map of this curve would be given in the same way. Therefore, it would be interesting to clarify the relation between the Weyl group expression and the differential operators.

Recently, the authors in [55] provides the analytic results on black hole perturbation theory from the quantization conditions. They consider the quantization conditions for A-periods, not B-periods. Therefore, it would be interesting to clarify the physical implications of this quantization conditions in the integrable systems or topological strings.

Acknowledgements

We would like to thank Sheldon Katz, Albrecht Klemm for helpful discussions and/or stimulating collaborations on related papers. Some of the computation based on mathematica were carried out on the computer sushiki at Yukawa Institute for Theoretical Physics in Kyoto University. The work of MH and YS was supported by the national Natural Science Foundation of China (Grants No.11675167 and No.11947301).

A An eigenvalue formula

Suppose $S$ is a real symmetric $2n \times 2n$ matrix, and $M$ is a real symplectic $2n \times 2n$ matrix that diagonalizes the symmetric matrix, i.e. we have

$$
\Sigma = \begin{pmatrix} 0 & I_n \\ -I_n & 0 \end{pmatrix}, \quad M^T \Sigma M = \Sigma, \quad M^T SM = \begin{pmatrix} C & 0 \\ 0 & D \end{pmatrix}, \quad \text{ (A.1)}
$$

where $C = \text{diag}\{c_1, c_2, \cdots, c_n\}, D = \text{diag}\{d_1, d_2, \cdots, d_n\}$ are real $n \times n$ diagonal matrices. Then we can show that the characteristic polynomial of the matrix $S\Sigma$ (or $\Sigma S$) is

$$
\det(S\Sigma - \lambda I) = \prod_{k=1}^{n}(\lambda^2 + c_k d_k).
\text{ (A.2)}
$$

So the eigenvalues of $S\Sigma$ are $\pm i\sqrt{c_k d_k}, k = 1, 2, \cdots n$. In the context of our physics problem, the two diagonal matrices are identical $C = D$, therefore the diagonal elements are completely determined by the symmetric matrix $S$, are thus independent of the choice of the symplectic matrix $M$.

The calculations are straightforward. Noticing $\Sigma^2 = -I$ and $(-\Sigma M^T)(\Sigma M) = I$, so the characteristic polynomial is

$$
\det(S\Sigma - \lambda I) = \det(-\Sigma M^T S\Sigma^2 M - \lambda I)
= \det\left( \begin{pmatrix} 0 & D \\ -C & 0 \end{pmatrix} - \lambda I \right).
\text{ (A.3)}
$$
It is now simple to verify the determinant is indeed the polynomial in the right hand side of (A.2).

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