Reviewer 2

• I expect the authors to rewrite the algorithm section. Firstly, the overview should be more neat and concise. Each notation, function, and equation should be clearly defined, and including a notation table would be helpful. Currently, the algorithm is presented in separate sections, with the basic concepts in Section 3 and the key technical contributions in Section 4, making it hard to follow.

A detail of the algorithm explaining the steps with their symbolic notation has been included in the supplementary file primarily due to space limitations in the main paper.

The algorithm stage is poorly organized. The writing needs to be improved.

The algorithm is reorganized in way that the contribution section is highlighted and rest of the explanations that describes the steps of the pipeline is moved to the supplementary file.

Reviewer 3

• The title of the paper does not accurately describe the content. The title describes plant phenotyping, while the dataset used in the evaluation section of the paper only contains maize and tomato, which are crops, and other plant types such as trees are not validated in the evaluation.

Maize and tomato dataset is used for benchmarking plant phenotyping in many existing works. Our work also depends on such dataset for determining the quality of the registration. Tomato is considered as a type of plant.

 The abstract of the paper states that the use of iterative beam search for correspondence matching is advantageous for handling large graphs. However, in the latter part of the paper, neither a definition of "large graphs" nor an experimental comparison and analysis of the paper's method for graphs of different scales is provided.

A description of the relative meaning of the large graph is added in Section 3. The large graph signifies the amount of nodes that can be handled during the generation of transition and emission cost. Viterbi algorithm tends to be computationally expensive for a graph with nodes around 50 whereas using the beam search, we can handle graph with nodes numbering 100-120.

• The reason for adopting the L1-medial skeletonization algorithm stated in paper 3.1 is that no significant advantages in adopting methods that introduce additional overhead and require segmentation approach.

However, the paper lacks a detailed explanation of the analytical process underlying this argument.

The process of skeletonization adopted by Chebrolu et al. involves multiple steps including segmentation using a machine learning algorithm, clustering to identify individual leaves or stems as distinct entities. Subsequently, they determined keypoints or skeleton nodes for each organ using self-organizing maps. Our analysis suggests that there are no distinct advantages of such strategies and hence we adopted L1-medial that is robust to complex dataset and effectively generates skeleton for the dataset used in our approach.

 The paper lacks a clear justification for the use of the Schunck et al. study as a dataset.

The dataset has been widely used for benchmarking and the same is quoted in the section 4.1 of the main paper. The choice of the dataset is attributed mainly because of its diversity and usage across different research papers focusing on plant phenotyping

• In paper 3.4, there is no explanation as to why the unknown parameters per node parameter for 3D affine transformation is 12.

An explanation is added to the supplementary file elaborating that the 12 parameters include 3X3 rotation matrix and 3X1 translation vector.

• In addition to the comparison with the method of Chebrolu et al., it would be beneficial to compare the paper with the method of the last two years to enhance the persuasive effect of the article.

-----to be answered-----

• The paper's depiction of the results is relatively clear, although it lacks a visual representation of the overall flow of the method.

An algorithmic pseudocode is added to the supplementary file (Section 1) to space constraints in the main paper. The pseudocode clearly emphasizes on the major steps included in the algorithm with respective input and corresponding output.

Reviewer 4

 The proposed method, while effective, appears to be computationally intensive and may require significant expertise to implement and fine-tune. This could limit its accessibility and practicality for broader applications without further simplification or optimization.

The runtime comparison shows that our method consumes little time while performing different task in the overall pipeline. The phenotyping is performed generally in a greenhouse environment therefore such timings can still be used without being a limitation.

• The experimental validation, although comprehensive, is limited to maize and tomato plants. It would be beneficial to evaluate the method on a more diverse set of plant species to demonstrate its generalizability and robustness.

The dataset used for comparison in our method is widely adopted across multiple research publications on phenotyping. It focuses on maize and tomato demonstrating complex growth and suitable for validating the result. On the other hand, another dataset for comparative analysis would be a surplus.

 While the paper compares the proposed method with a few existing techniques, a more extensive comparison with a broader range of state-of-the-art methods would strengthen the validation.

-----to be answered-----

 Explore potential optimizations to reduce the computational complexity and make the method more accessible for practical applications.

The runtime comparison shows that our method consumes little time while performing different task in the overall pipeline. The phenotyping is performed generally in a greenhouse environment therefore such timings can still be used without being a limitation.