## FASTLINK

**RRID:** SCR_009177  
**Type:** Tool

### Proper Citation

FASTLINK (RRID:SCR_009177)

### Resource Information

<table>
<thead>
<tr>
<th></th>
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</thead>
<tbody>
<tr>
<td><strong>Description:</strong></td>
<td>Software application (entry from Genetic Analysis Software)</td>
</tr>
<tr>
<td><strong>Resource Name:</strong></td>
<td>FASTLINK</td>
</tr>
<tr>
<td><strong>Proper Citation:</strong></td>
<td>FASTLINK (RRID:SCR_009177)</td>
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<tr>
<td><strong>Resource Type:</strong></td>
<td>Resource, software resource, software application</td>
</tr>
<tr>
<td><strong>Keywords:</strong></td>
<td>gene, genetic, genomic, c, unix, vms, ms-dos, .. and can also run in parallel on shared memory unix machines</td>
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<td><strong>Resource ID:</strong></td>
<td>SCR_009177</td>
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<tr>
<td><strong>Website Status:</strong></td>
<td>Last checked up</td>
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<tr>
<td><strong>Alternate IDs:</strong></td>
<td>nlx_154309</td>
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<tr>
<td><strong>Abbreviations:</strong></td>
<td>FASTLINK</td>
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<tr>
<td><strong>Mentions Count:</strong></td>
<td>39</td>
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</table>

### Ratings and Alerts

No rating or validation information has been found for FASTLINK.

No alerts have been found for FASTLINK.
Usage and Citation Metrics

We found 39 mentions in open access literature.

**Listed below are recent publications.** The full list is available at [FDI Lab - SciCrunch Infrastructure](https://fdi.lab).


Ullah I, et al. (2016) Pathogenic mutations in TULP1 responsible for retinitis pigmentosa identified in consanguineous familial cases. Molecular vision, 22, 797-815.

Khan SY, et al. (2016) FOXE3 contributes to Peters anomaly through transcriptional


