

## BAB VIII

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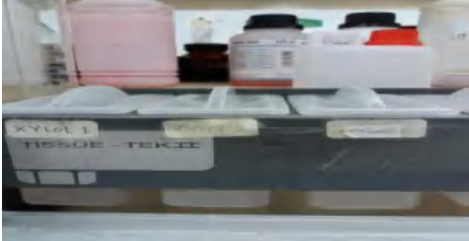


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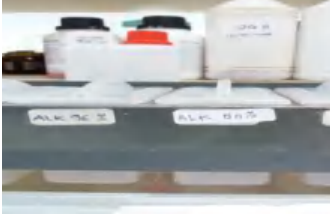








LAMPIRAN 1

1. Prosedur pewarnaan jaringan biologis penderita kanker payudara

<p>1. Potong blok paraffin dengan mikrotom pada ketebalan 3-4<math>\mu</math></p> 	<p>2. Celupkan kedalam Waterbath</p> 
<p>3. Ambil potongan jaringan dengan slide lalu tiriskan</p> 	<p>4. Tulis pada slide kode sesuai blok paraffin dengan pensil</p> 
<p>5. Panaskan slide diatas Hot Plate selama 1 jam</p> 	<p>6. Dinginkan slide lalu masukkan kedalam keranjang slide</p> <p>7. Deparafinasi ( Xylol I, Xylol II, Xylol III) masing-masing 5 menit</p> 











<p>8. Rehidrasi (Alkohol 96%, Alkohol 80%, Alkohol 70%), masing-masing selama 5 menit</p> 	<p>9. Cuci air mengalir selama 5 menit</p> 
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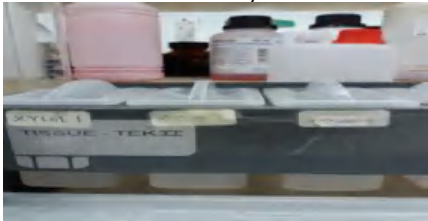
<p>10. Angkat dari air lalu masukkan slide kedalam larutan Blocking Endogen Peroksida, rendam selama 15 menit</p> 	<p>11. Cuci air mengalir selama 5 menit</p> 
<p>12. Masukkan keranjang berisi slide kedalam decloaking yang berisi larutan Antigen Retrieval Decloaking Chamber, lalu letakkan slide pada rack holder</p> 	<p>13. Masukkan rack holder kedalam decloaking, lalu tutup</p> 
<p>14. Atur waktu yaitu selama 40 menit pada suhu 95 derajat.</p> 	<p>15. Dinginkan, dengan mengeluarkan slide dari decloaking dan disimpan pada suhu ruangan</p> <p>16. Setelah dingin, cuci dalam larutan PBS 2x masing-masing selama 5 menit</p> <p>17. Tandai slide dengan memberi lingkaran sekitar jaringan</p>





	
<p>18. Atur slide pada baki slide</p> 	<p>19. Ambil satu per satu lalu tetesi Background Sniper lalu inkubasi selama 30 menit</p> 

<p>20. Buang larutan background sniper dengan cara ditiriskan pada tisu</p> <p>21. Tetesi Antibody Primer lalu inkubasi selama 1 jam pada suhu ruang</p> <p>22. Cuci PBS 2x selama 5 menit</p> 	<p>23. Tetesi Trekkie Universal lalu diamkan selama 30 menit</p> 
<p>24. Tiriskan pada tissue lalu tetesi Trekkavidin-HRP lalu diamkan selama 30 menit</p> 	<p>25. Cuci PBS 2x dengan cara merendam slide selama 5 menit</p> 
 <p>ambil menunggu pencucian, at larutan DAB dengan cara campurkan cromogen DAB 1</p>	<p>27. Amati jaringan jika sudah menunjukkan warna coklat, langsung direndam ke dalam air selama 5 menit</p>

<p>tetes + Substrat buffer 1 ml (dicampur dalam tabung bersih) lalu tetesi ke jaringan.</p> 	
<p>28. Rendam dengan hematoxylin Meyer 5 menit</p> 	<p>29. Cuci air mengalir selama 5 menit</p> 
<p>30. Dehidrasi ( Alkohol 70%, Alkohol 80%, Alkohol 96%) masing-masing 5 menit</p> 	<p>31. Clearing ( Xylol I, Xylol II, Xylol III)</p> 
<p>32. Keringkan slide lalu tetesi dengan entelan lalu tutup dengan deck glass.</p>	<p>33. Amati di mikroskop</p>



