# **Electrophoresis, Blotting and Immunodetection**

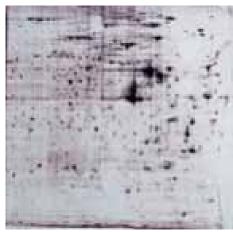
## Hybridisation equipment, blotting membranes



Standard immunodataction mathod



Rapid immunodetection method. Standard vs immunodetection methods using BCIP substrate as Immobilon™-P transfer membranes



2-D electrotransfer onto Immobilon™-P using silver stain method



## Filters, transfer membranes, PVDF, blotting, Immobilon™



#### Immobilon™-l

- · For general immunodetection and amino acid analysis
- Rapid immunodetection method reduces detection time by two hours
- · High mechanical strength for easy handling

### Immobilon™-FL

- The first transfer membrane optimised for fluorescence applications
- Extremely low background
- Compatible with all commonly used fluorescent probes
- Can be used at all excitation and emission wavelengths
- · Ideal for multiplexing

High protein binding 0.45µm membrane with low backgrounds. Eliminates blocking and subsequent wash steps in Western blots with new rapid immunodetection protocol that does not compromise specificity or sensitivity. Suitable for chromogenic chemiluminescent, chemifluorescent and radioactive detection. Transfer proteins from a variety of gel matrices by electroelution. Immobilon™-P offers increased solvent and staining capabilities and has a higher signal-to-noise ratio than does nitrocellulose. High mechanical tensile strength allows easy handling without cracking or breaking.

Technical Specification - Specific

|                              | Immobilon™-P   | lmmobilon™-FL  |  |
|------------------------------|--|--|--|
| Description                  | The original PVDF membrane for Western, dot and other protein blots  | Optimised for fluorescent<br>Immunodetection applications      |  |
| Material                     | Hydrophobic PVDF   | Hydrophobic PVDF   |  |
| Pore size, µm                | 0.45   | 0.45   |  |
| Applications                 | Western blotting, binding assays,<br>amino acid analysis, N-terminal protein<br>sequencing, dot/slot blotting,<br>glycoprotein visualisation,<br>lipopolysaccharide analysis | Western blotting, dot/slot blotting                            |  |
| Detection                    | Chemiluminescent, chromogenic, radioactive   | Fluorescent, chemifluorescent, chromogenic, chemiluminescent   |  |
| Protein binding capacity, mg | Insulin: 85<br>BSA: 131<br>Goat IgG: 294   | Insulin: 85<br>BSA: 131<br>Goat IgG: 131                       |  |
| Compatible stains            | Coomassie® Brilliant Blue, amido black,<br>india ink, Ponceau-S red, collodial gold,<br>GPTS, toluidine blue, transillumination,<br>Sypro® Ruby                              | Coomassie® Brilliant Blue, amido black,<br>Ponceau-S red, GPTS |  |

All proteins may not behave the same on a membrane surface; variability in properties such as charge, density, conformation, or hydrophobicity may necessitate use of another Immobilon™ PVDF membrane for Western blotting.

## 0.45µm Immobilon™-P

| Catalogue No | Alt. No   | Format | Dimensions, mm | Pack qty |
|--------------|-----------|--------|----------------|----------|
| FDR-520-027U | IPVH08100 | Sheets | 80 x 100       | 10       |
| FDR-520-050C | IPVH15150 | Sheets | 150 x 150      | 10       |
| FDR-520-060W | IPVH20200 | Sheets | 200 x 200      | 10       |
| FDR-520-080Q | IPVH00010 | Roll   | 265 x 3,750    | 1        |

### 0.45µm Immobilon™-FL

| Catalogue No | Alt. No   | Format | Dimensions, mm | Pack qty |
|--------------|-----------|--------|----------------|----------|
| FDR-523-010T | IPFL10100 | Sheets | 100 x 100      | 10       |
| FDR-523-020Q | IPFL20200 | Sheets | 200 x 200      | 10       |
| FDR-523-030N | IPFL00010 | Roll   | 265 x 3,750    | 1        |

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